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(54) **MOLECULAR ORIGIN OF ALLERGY**

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CPC **A23C 9/1465** (2013.01); **A23C 9/20** (2013.01); **A23J 3/00** (2013.01); **A23J 3/08** (2013.01); **A23J 3/14** (2013.01); **A23K 10/20** (2016.05); **A23L 5/27** (2016.08); **A23L 5/273** (2016.08); **B01D 15/362** (2013.01); **B01D 15/3809** (2013.01); **C07K 1/18** (2013.01); **C07K 1/22** (2013.01); **C07K 14/415** (2013.01); **G01N 33/6854** (2013.01); **A23V 2002/00** (2013.01); **A23V 2200/03** (2013.01); **A23V 2300/30** (2013.01); **G01N 2800/24** (2013.01); **G01N 2800/50** (2013.01)

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See application file for complete search history.

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(57) **ABSTRACT**

The present invention relates to compositions and methods for modulating or detecting allergy in a subject. The invention may be used to reduce allergenicity of compositions, such as food products, or to stimulate immunogenicity of products, such as vaccines by removal of cationic proteins resulting from transcription infidelity. The invention may be used in any mammal such as human.

18 Claims, 11 Drawing Sheets

(4 of 11 Drawing Sheet(s) Filed in Color)

Specification includes a Sequence Listing.

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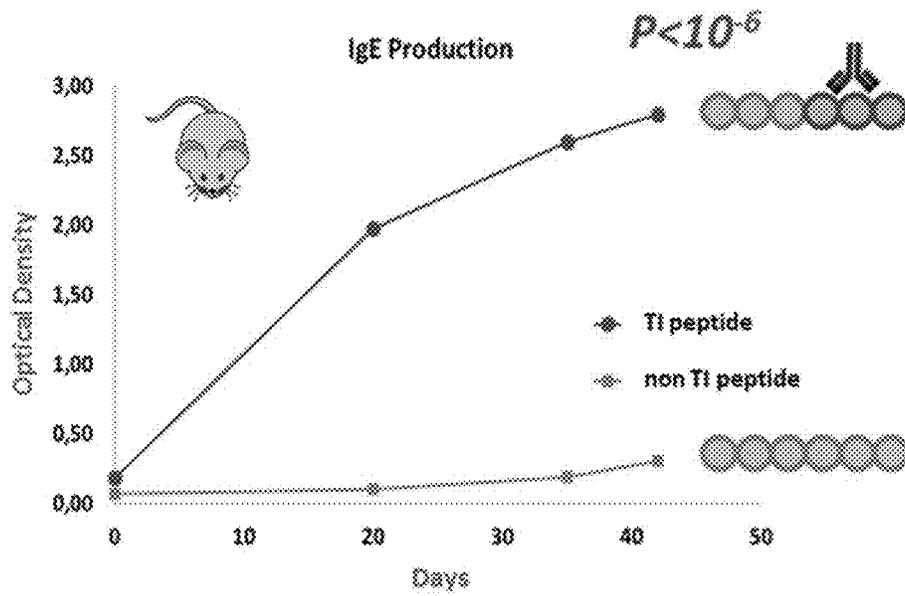
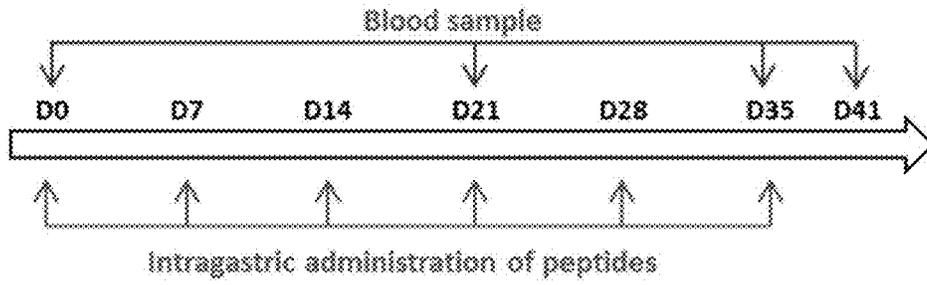


Figure 1

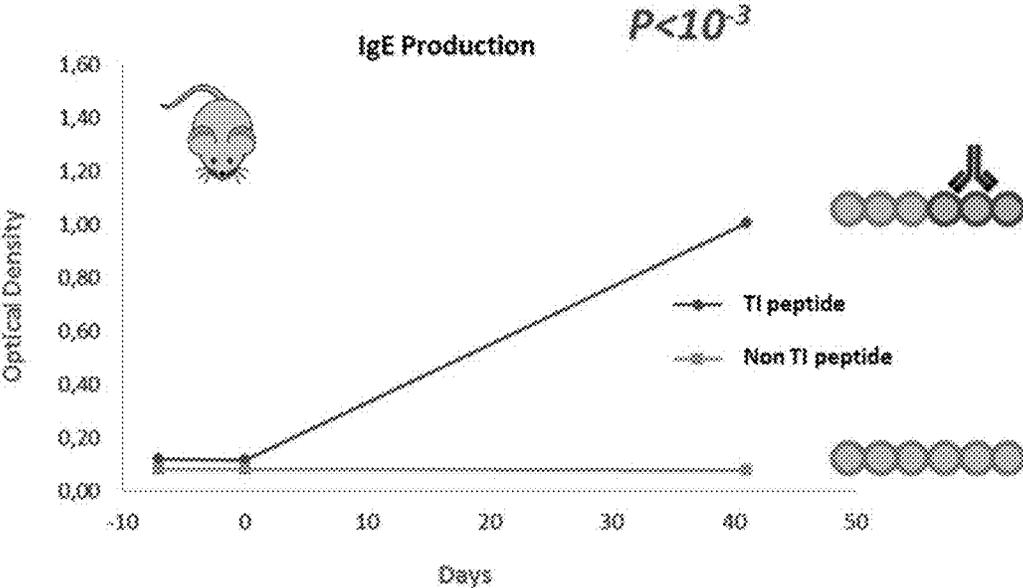
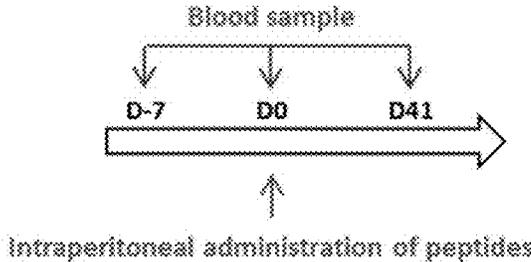


Figure 2

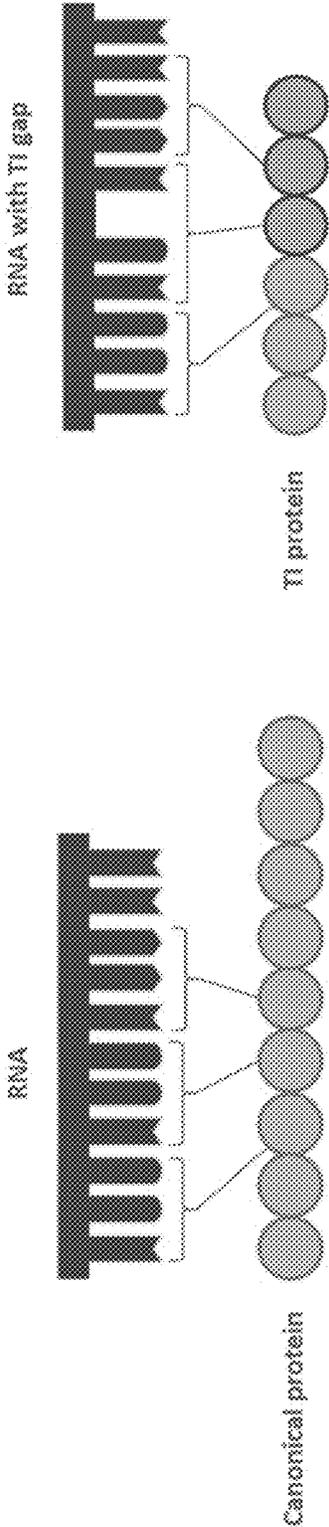


Figure 3

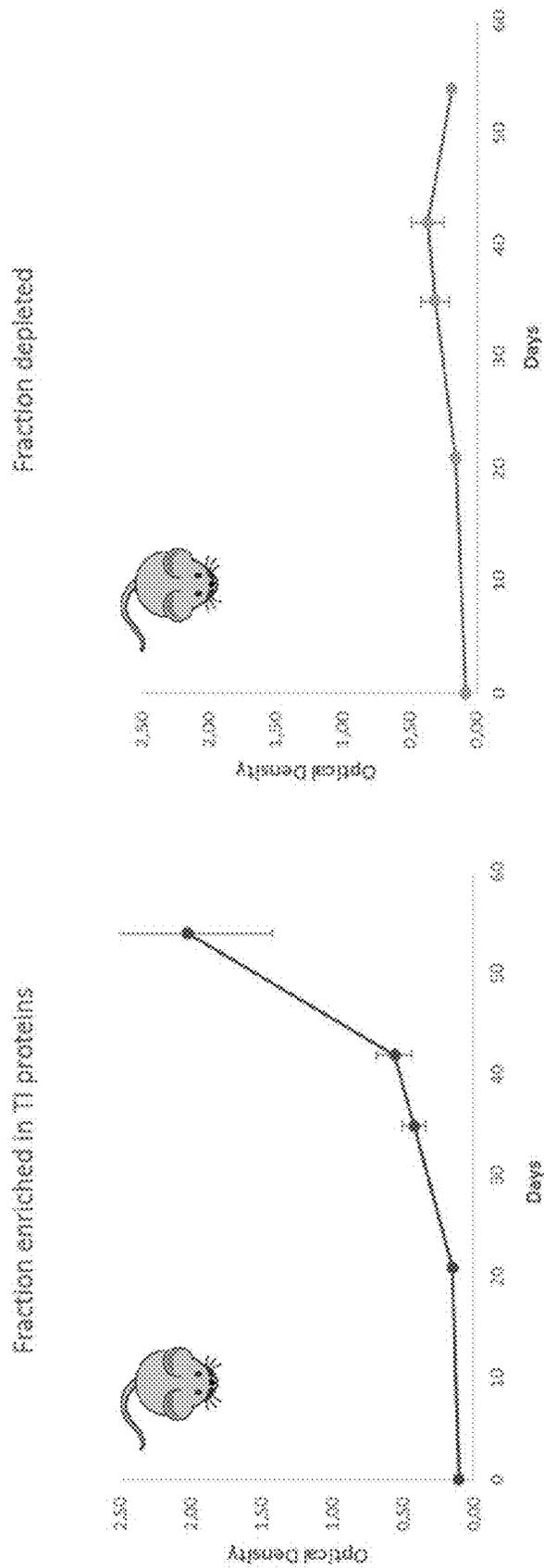


Figure 4

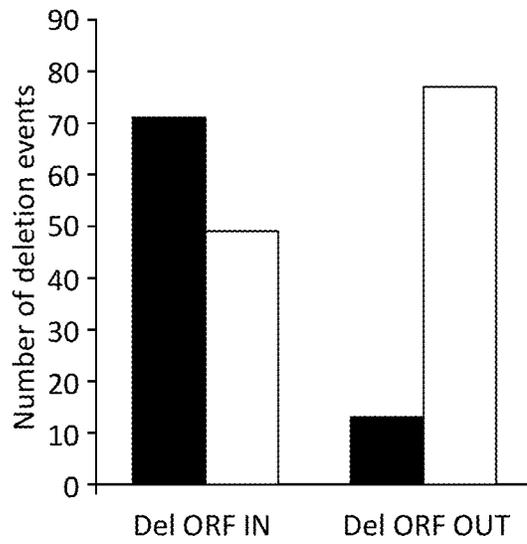


Figure 5

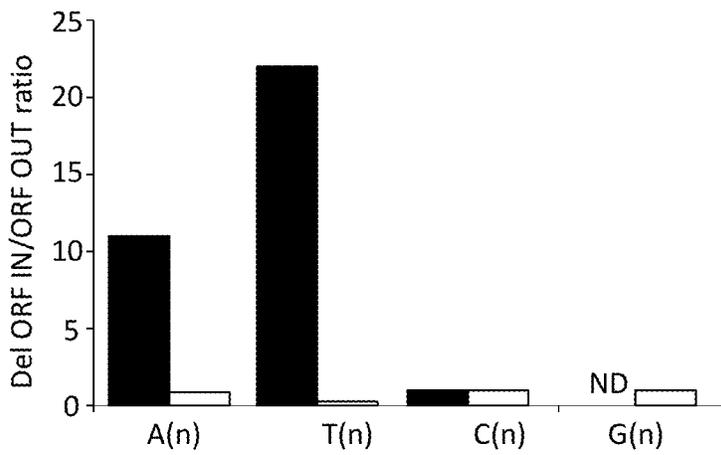


Figure 6

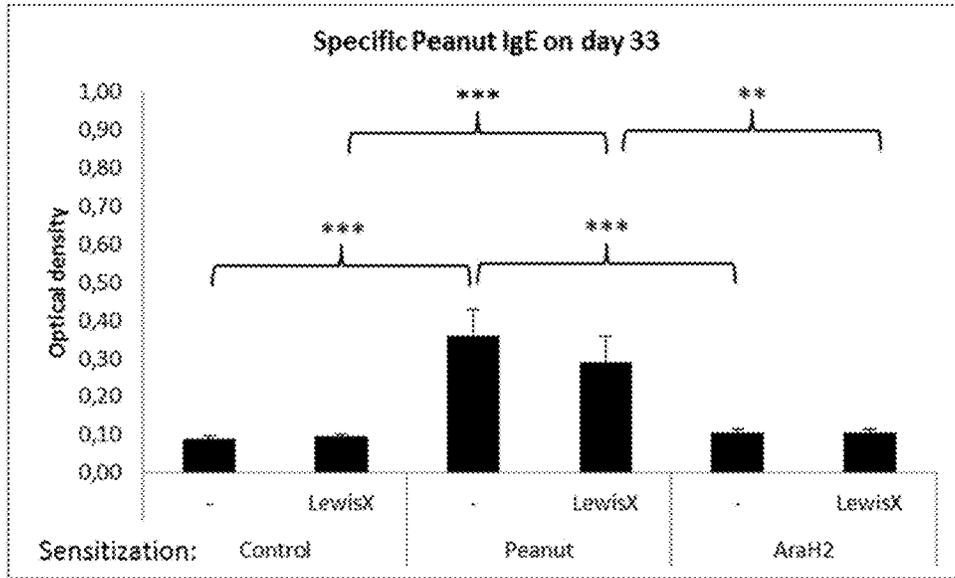


Figure 7A

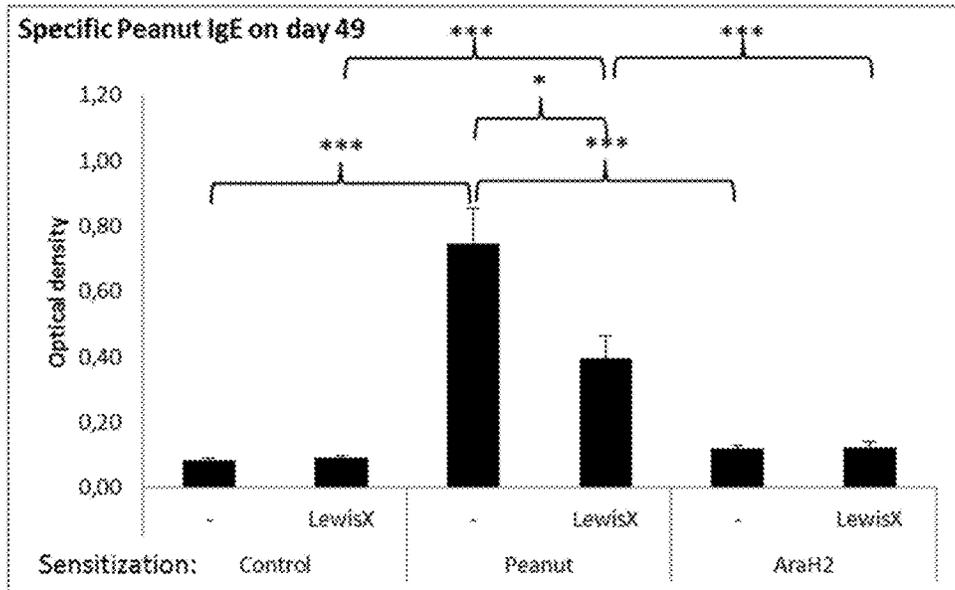
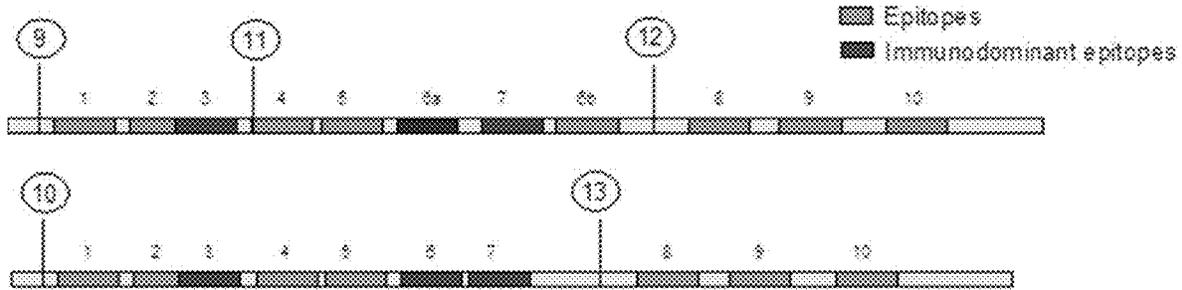


Figure 7B



11

 epitope 3 epitope 4 epitope 5 epitope 6 epitope 7

 ref DRRCQSGLEBANLRFCEQHLMKIQDDESYGRDPYSPSQDFYSPSQDFDRRDYSPSPYDRRGAGSSQHQERCNE

 30 DRRCQSGLEBANLRFCEQHLMKIKKSNVTRIHMGGTKTALVRIKRTALVTRTDVIRKALVHMIGEALDLLSTKRGVAMS

 31 DRRCQSGLEBANLRFCEQHLMKIQDDESYGRDPYSLVRIKRTALVTRTDVIRKALVHMIGEALDLLSTKRGVAMS

 32 DRRCQSGLEBANLRFCEQHLMKIQDDESYGRDPYSPFERIKRTALVTRTDVIRKALVHMIGEALDLLSTKRGVAMS

 33 DRRCQSGLEBANLRFCEQHLMKIQDDESYGRDPYSPSQDFYSPSQDFDRRDYSLVHMIGEALDLLSTKRGVAMS

 34 DRRCQSGLEBANLRFCEQHLMKIQDDESYGRDPYSPSQDFYSPSQDFDRRDYSPSPYDRRGAGSLSTKRGVAMS

ID	Antigen	Group	Pos	Ref pt	Ref seq (Da)	Ref len	T1 prot pt	T1 prot seq	Len	T1 prot pt	No seq	Cov	No seq	Len	Epitope
30	Ara h 2	stop_319	51	6.23	20114.26	172	11.97	11470.13	106	12.41	1	983	1	16	epitope_4 stop_1
31	Ara h 2	stop_319	67	6.23	20114.26	172	10.30	12046.98	106	12.21	2	796	1	16	epitope_6 stop_1
32	Ara h 2	stop_319	69	6.23	20114.26	172	10.30	13057.98	106	12.21	1	732	1	2	epitope_7 start_1
33	Ara h 2	stop_319	86	6.23	20114.26	172	6.70	12082.66	106	7.00	1	331	1	17	epitope_7 stop_10
34	Ara h 2	stop_319	97	6.23	20114.26	172	8.48	12034.46	106	11.62	8	7995	1	11	epitope_7 stop_21

12

 ref GAGSSQHQERCCELNEFENNQRCEALQQ

 35 GAGSSQHQERCCELNEFENNKGACARHCNR

 36 GAGSSQHQERCCELNEFENNQGACARHCNR

ID	Antigen	Group	Pos	Ref pt	Ref seq (Da)	Ref len	T1 Prot pt	T1 prot seq	Len	T1 prot pt	No seq	Cov	No seq	Len	Epitope
35	Ara h 2	stop_370	114	6.23	20114.26	172	7.21	14176.85	123	9.81	28	14975	1		epitope_8 start_10
36	Ara h 2	stop_370	115	6.23	20114.26	172	6.82	14176.81	123	8.57	40	33373	4	1	epitope_8 start_18

13

 epitope 8 epitope 9 epitope 10

 37 GAGSSQHQERCCELNEFENNQRCEALQQIMENQSDRIQGGQSQDFEELRLPQQQLGHSVATWTSKVAETDTRHLQKKKKRKKIAYI

 38 GAGSSQHQERCCELNEFENNKGACARHCNRNRSWETHAIGCRGGRNENSSRGS SGTCLNSAALGHSVATWTSKVAETDTRHLQKKKKRKKIAYI

 39 GAGSSQHQERCCELNEFENNQRCEALQQIMENQSDRIQGGQSQDFEELRLPQQQLGHSVATWTSKVAETDTRHLQKKKKRKKIAYI

 40 GAGSSQHQERCCELNEFENNQRCEALQQIMENQSDRIQGGQSQDFEELRLPQQQLGHSVATWTSKVAETDTRHLQKKKKRKKIAYI

ID	Antigen	Group	Pos	Ref pt	Ref seq (Da)	Ref len	T1 Prot pt	T1 prot seq	Len	T1 prot pt	No seq	Cov	No seq	Len	Epitope
37	Ara h 2	stop_474	102	6.23	18757.85	160	9.74	15968.54	177	11.45	44	22303	1		epitope_8 start_18
38	Ara h 2	stop_474	103	6.23	18757.85	160	8.90	15968.30	177	11.41	30	24031	2	1	epitope_8 start_17
39	Ara h 2	stop_474	129	6.23	18757.85	160	8.16	20262.54	177	10.60	2	1190	1	28	epitope_9 start_1
40	Ara h 2	stop_474	145	6.23	18757.85	160	8.59	20704.28	177	10.62	15	10708	2	18	epitope_10 start_1

Figure 8

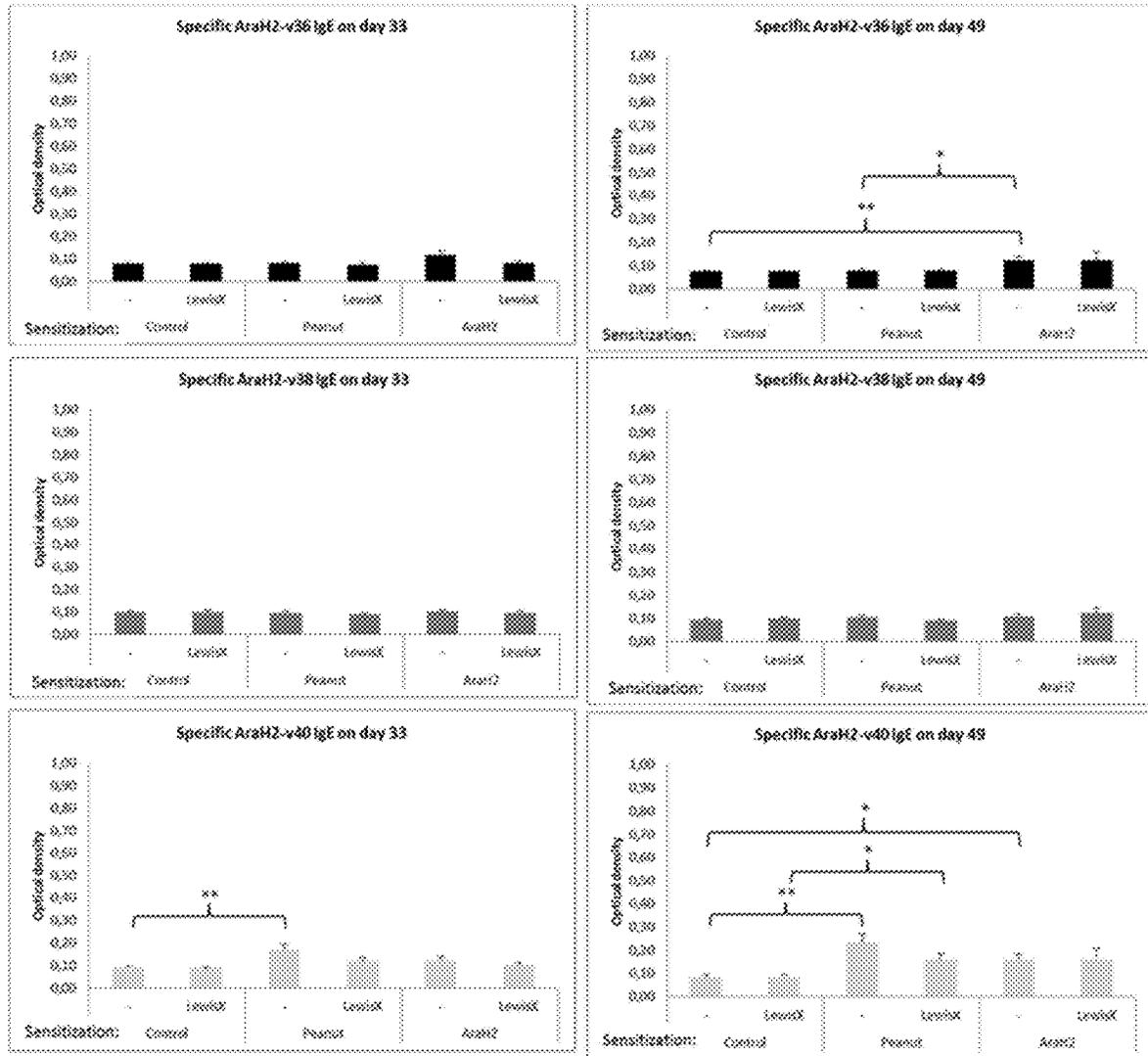


Figure 9

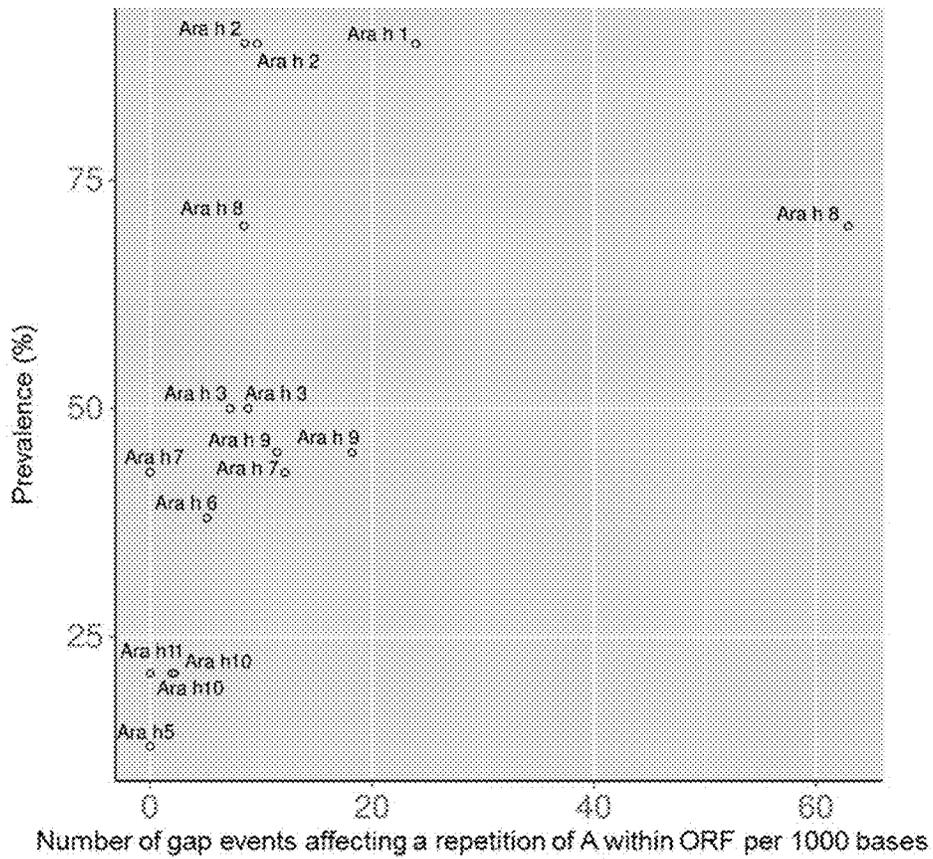


Figure 10

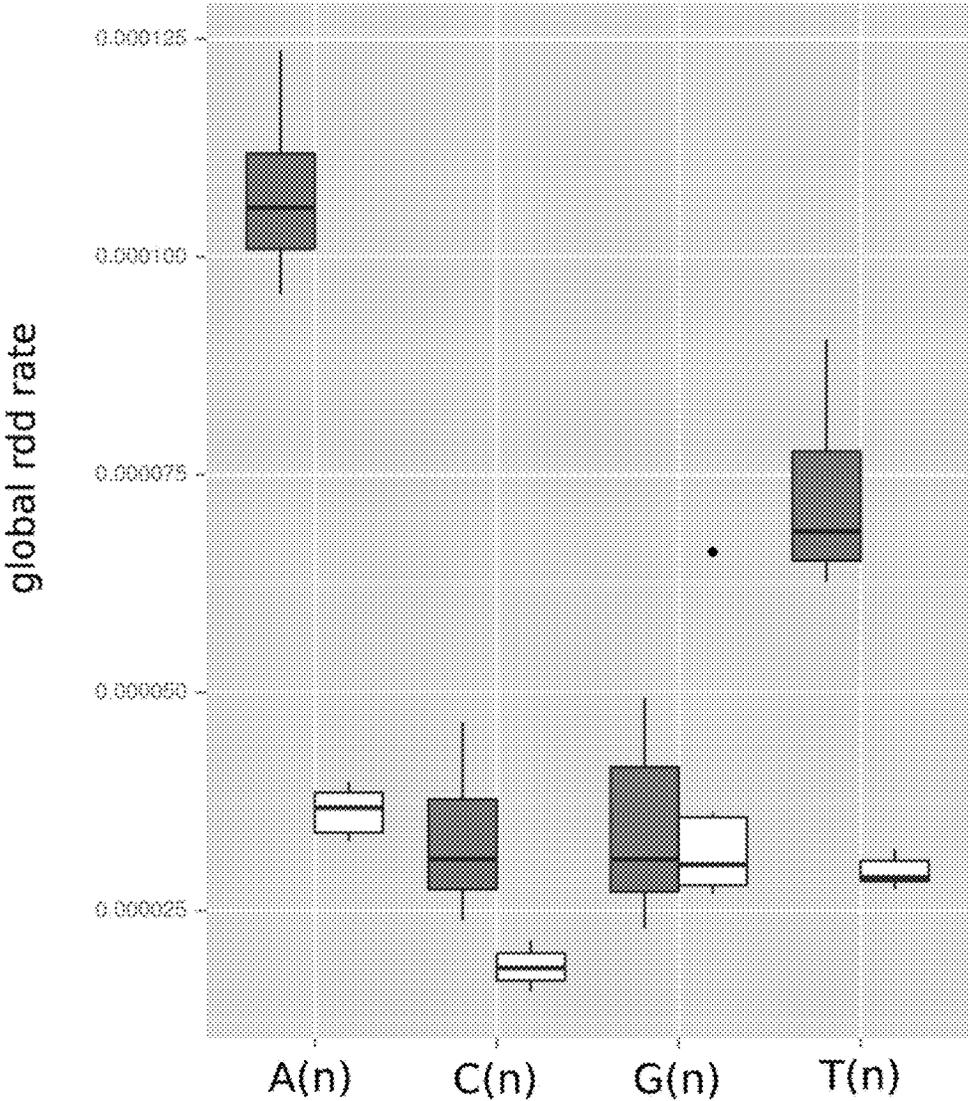


Figure 11

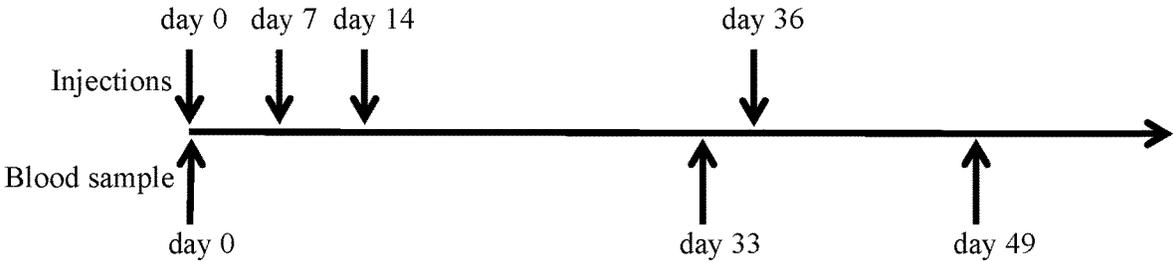


Figure 12

MOLECULAR ORIGIN OF ALLERGY**CROSS-REFERENCE TO RELATED APPLICATION**

This application is the U.S. national stage application of International Patent Application No. PCT/EP2017/056576, filed Mar. 20, 2017.

The Sequence Listing for this application is labeled "Seq-List.txt" which was created on Sep. 13, 2018 and is 199 KB. The entire content of the sequence listing is incorporated herein by reference in its entirety.

The present invention relates to compositions and methods for modulating or detecting allergy in a subject. The invention stems from the discovery, by the inventors, of a molecular origin of allergy in mammals and has wide uses in medical, nutritional, cosmetic or agricultural industries, for instance. The invention may be used to reduce allergenicity of compositions, such as food products, or to stimulate immunogenicity of products, such as vaccines. The invention may be used in any mammal such as human.

BACKGROUND

Clinical manifestations of allergic reactions are highly diversified affecting virtually all organs that can be in contact with external environment i.e. respiratory, skin, digestive and geneco urinary and can culminate in systemic anaphylactic shock with more or less serious haemodynamic consequences. IgE antibody are the cornerstone of Type I hypersensitivity causing diseases such as Asthma, Hay Fever, Eczema, Urticaria, food allergies and anaphylaxis. IgE are both circulating and anchored to FcεRI expressed at the surface of mast cells, basophils, eosinophils, monocytes, macrophages and

Langerhans cells. Allergen induced cross-linking of IgE anchored to FcεRI on tissue mast cells elicit a cascade of events liberating histamine and multiple cytokines that contribute to both amplification of the reaction by recruiting circulating basophils also armed with IgE anchored to FcεRI. While these events have been characterized in great details, the simple question as to why any given protein that for most individuals is harmless becomes a potentially lethal allergen for others remains unanswered. The work presented here explains that the initial trigger of IgE production stems not from known allergens but from minute amount of variants translated from mRNA with frame shift caused by transcriptional errors.

SUMMARY

The invention stems from the discovery, by the inventors, of a molecular origin of allergy in mammals. The invention thus provides novel compositions and methods for detecting, controlling or modulating an immune response or allergy in mammals. The invention particularly stems from the discovery, by the inventors, that allergy is triggered by proteins (or epitopes thereof) generated by transcription infidelity ("TI") in mammals. In particular, as previously demonstrated by the inventors, TI generates aberrant proteins with modified C-terminal ends. Continuing their investigations, the inventors have now surprisingly found that TI gaps create proteins which acquire immunogenicity in mammals and trigger allergy in vivo. The inventors have also found that such proteins resulting from TI gaps are essentially cationic and that removing such proteins from food or other compositions generates hypoallergenic compositions.

The present invention thus provides novel compositions and methods for detecting, monitoring and modulating immunogenicity and allergy in mammals.

An object of the invention more particularly resides in a method for reducing allergenicity or immunogenicity of a composition, the method comprising treating the composition to remove cationic proteins.

An object of the invention resides in a method for reducing allergenicity or immunogenicity of a composition, the method comprising treating the composition to remove proteins resulting from transcription infidelity, more particularly proteins having a sequence resulting from a transcription infidelity gap.

The composition may be any composition such as a food, feed, pharmaceutical product, veterinary product, cosmetic product, etc.

In a particular embodiment, the invention provides a method for preparing a food product comprising (i) providing a food product preparation, (ii) treating the food product preparation to remove cationic proteins therefrom and (iii) optionally formulating the treated food product with one or more suitable excipients.

Another particular embodiment of the invention relates to a method for preparing a pharmaceutical product comprising (i) providing a pharmaceutical product preparation, (ii) treating the pharmaceutical product preparation to remove cationic proteins therefrom and (iii) optionally formulating the treated pharmaceutical product with one or more suitable excipients.

The invention also provides food compositions comprising a food product and a suitable excipient, wherein the food product contains less than 1% by weight of cationic proteins, more preferably less than 0.5%, less than 0.3%, less than 0.2%, or less than 0.1%.

The invention also provides pharmaceutical compositions comprising a pharmaceutical/veterinary product and a suitable excipient, wherein the pharmaceutical/veterinary product has been treated to contains less than 1% by weight of cationic proteins, more preferably less than 0.5%, less than 0.3%, less than 0.2%, or less than 0.1%.

The invention also relates to a method of treating a subject, comprising administering to the subject an effective amount of a pharmaceutical or veterinary product as defined above.

The invention also provides a method for detecting a subject having predisposition to allergy, comprising measuring in a sample from said subject the level of IgE directed against proteins having a sequence resulting from a transcription infidelity, wherein a difference in said level as compared to a control value indicates a subject having predisposition to allergy.

A further object of the invention is a cationic protein or peptide having a sequence resulting from transcription infidelity, for use as an adjuvant (e.g., to stimulate antibody production in a mammal).

The invention also provides a method of inducing or stimulating antibody production in a mammal, comprising administering to the mammal a cationic protein or peptide having a sequence resulting from transcription infidelity.

The invention further provides a method for producing IgE comprising (i) administering to a non-human mammal a cationic protein or peptide resulting from transcription infidelity under conditions allowing induction of IgE production, and (ii) collecting IgE produced.

The invention further provides a method for producing IgE comprising (i) administering to a non-human mammal a cationic protein or peptide resulting from transcription infidelity

delity under conditions allowing induction of IgE production, (ii) collecting IgE-producing cells and (iii) deriving monoclonal and/or humanized IgE from said collected cells. Step (iii) typically comprises producing hybridoma, clonal selection of hybridomas and production of monoclonal antibodies.

The invention also concerns a method of making anti-IgE antibodies, comprising (i) administering to a non-human mammal a cationic protein or peptide resulting from transcription infidelity under conditions allowing induction of antibodies, (ii) collecting antibodies produced and (iii) selecting antibodies that bind Fc receptor.

A further object of the invention is a pharmaceutical composition comprising an antibody that binds Fc receptor.

The invention also provides a vaccine composition comprising an immunogen and a protein or peptide having a sequence resulting from transcription infidelity.

LEGEND TO THE FIGURES

The patent or application file contains at least one drawing executed in color. Copies of this patent or patent application publication, with color drawing(s), will be provided by the Office upon request and payment of the necessary fee.

FIG. 1. IgE production after intragastric administration of TI and non-TI peptides. The days of administration of the peptides and the collection of blood from the mice are indicated in the upper part. The assay of IgEs in the blood is carried out by the ELISA technique and the results are expressed as optical density. The difference between the two peptides is significant ($p < 10^{-6}$).

FIG. 2. IgE production after intraperitoneal administration of TI and non-TI peptides. The days of administration of the peptides and the collection of blood from the mice are indicated in the upper part. The assay of IgEs in the blood is carried out by the ELISA technique and the results are expressed as optical density. The difference between the two peptides is significant ($p < 10^{-3}$).

FIG. 3. A TI gap (in red) on the RNA causes a shift in the reading frame of the protein. The sequence of the protein resulting from the gap is very different from the normal protein.

FIG. 4. Administration to mice of fractions enriched in and depleted of cationic proteins (6 mice per group). From blood samples, the assay of IgEs is carried out by the ELISA technique and the results are expressed as optical density. The error bars indicate the variability of the measurement for the whole group.

FIG. 5: Deletions are mainly located within ORF of allergens. Number of deletions has been determined for allergens (black) and non-allergens (white). The position of the deletion is then defined as ORF IN (located within the coding sequence) or ORF OUT (located within untranslated regions).

FIG. 6: Ratio between the number of deletions located within ORF and within UTR, for allergens (black) and non-allergens (white), for deletions affecting repetitions of A, T, C and G. ND: non determined (there is no deletion affecting repetitions of G in the allergens).

FIGS. 7A and 7B: IgE antibody responses to peanut on day 33 (A) and on day 49 (B). Mice (n=10 per treatment group, per experiment) received peanut extract (400 µg protein) or recombinant AraH2 (400 µg) by intra peritoneal injection on days 0, 7, 14 and 36, with or without LewisX adjuvant. Serum samples (days 33 and 49) were analyzed for specific IgE antibody by ELISA. Data are shown as mean

(±SEM). Non parametric Wilcoxon test are performed (* $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$).

FIG. 8: Positions of deletions affecting Ara h 2 and prediction of TI proteins and peptides. Two isoforms are shown. Epitopes from each isoform are represented in grey and immunodominant epitopes are represented in black. Vertical bars indicate the positions of group of deletions implying the same stop codon. For some groups, sequences are shown. Transcription infidelity (TI) sequences are bold; epitopes are underlined. Diagrams 11 (ref, residues 11-87 of SEQ ID NO: 126; 30, residues 11-87 of SEQ ID NO: 121; 31, residues 11-87 of SEQ ID NO: 122; 32, residues 11-87 of 123; 33, residues 11-87 of SEQ ID NO: 124; 34, residues 11-87 of SEQ ID NO: 125), 12 (ref, residues 62-92 of SEQ ID NO: 130; 35, residues 74-104 of SEQ ID NO: 126; 36, residues 74-104 of SEQ ID NO: 127) and 13 (top row, residues 74-153 of SEQ ID NO: 132; 37, residues 62-158 of SEQ ID NO: 128; 38, residues 62-158 of SEQ ID NO: 129; 39, residues 62-158 of SEQ ID NO: 130; 40, residues 62-158 of SEQ ID NO: 131) also contain biochemical and bioinformatical characteristics of these TI variants. Stop codon of reference sequence is labelled with an asterisk.

FIG. 9: IgE antibody responses to AraH2-v36 on day 33, on day 49; IgE antibody responses to AraH2-v38 on day 33, on day 49; IgE antibody responses to AraH2-v40 on day 33, on day 49. Mice (n=10 per treatment group, per experiment) received Peanut extract (400 µg protein) or recombinant AraH2 (400 µs) by intra peritoneal injection on days 0, 7, 14 and 36, with or without LewisX adjuvant. Serum samples (days 33 and 49) were analyzed for specific IgE antibody by ELISA. Data are shown as mean (±SEM). Non parametric Wilcoxon test are performed (* $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$).

FIG. 10: Correlation between the number of gap events in peanut Ara h 1-3 and Ara h 5-11 transcripts (said gap events affecting a repetition of A within ORF per 1000 bases) and prevalence of the peanut allergens.

FIG. 11: Comparison of global RDD rate computed on all transcript positions, for peanuts (dark grey) and green beans (white), for deletions affecting repetitions of A, C, G and T bases within the coding regions ORF.

FIG. 12: Diagram of immunization protocol

DETAILED DESCRIPTION OF THE INVENTION

The present invention provides an unified model that explains how any given protein variant, produced by translation of mRNA carrying specific transcription infidelity (TI) event, causes deregulation of natural immunoglobulin specific properties and is at the origin of allergy. More particularly, the present invention defines that known allergens translated from RNA transcribed by a canonical base pairing are unable to elicit modification of natural immunoglobulin properties. On the contrary, their transcription infidelity (TI) variants are clearly able to cause these modifications and trigger allergy.

The invention also provides methods for reducing allergenicity of various food or pharmaceutical compositions or products (such as milk or dairy products, peanut products, peanuts, pharmaceutical preparations, etc.). More particularly, the invention provides methods to remove and/or chemically modify allergy triggers in order to reduce their capacity to elicit change in natural immunoglobulin properties. The invention is exemplified for various forms of allergy (e.g., milk, peanut, mite allergy) and in various mammalian species including human.

Using a murine model of allergy, we surprisingly observed that a protein incapable of inducing IgE production acquired this capacity when it underwent a gap modification resulting from transcription infidelity. This suggests a molecular origin of allergy resulting not from classic allergens but from their protein variants resulting from transcription infidelity. To confirm this hypothesis, we sequenced the RNA of cow's milk and observed the presence of a gap near the main B epitopes of milk proteins such as α_{s1} -casein and β -lactoglobulin. We also demonstrated that gap-modified proteins resulting from transcription infidelity had a cationic nature. We thus proceeded to a fractionation of the whey so as to enrich it in TI-produced cationic proteins and then to a comparison of the ability of the enriched and depleted fractions to induce IgE. Remarkably, and unexpectedly, the enriched fraction administered at a dose of 57 μ g induces IgE production equivalent to that observed with 1.9 mg of whey protein. In contrast, 57 μ g of the depleted fraction does not induce significant IgE production. These results thus show that low abundant variants of proteins can trigger production of IgE which, by extension toward the N-ter portion, also bind the normal protein. The invention thus discloses a molecular mechanism at the origin of allergy and provides new approaches for early diagnosis, treatment and prevention of allergies, and for modulating the immune response.

An object of the invention more particularly resides in a method for reducing allergenicity or immunogenicity of a composition, the method comprising treating the composition to remove proteins resulting from transcription infidelity.

Another object of the invention resides in a method for reducing allergenicity or immunogenicity of a composition, the method comprising treating the composition to remove cationic proteins.

The invention also relates to compositions obtainable by the above methods, as well as the uses thereof.

Within the context of the present invention, the term "immunogenicity" designates for instance the ability of a composition or protein or molecule to induce an immune response in a mammal. This includes for instance the ability to trigger antibody production or a T cell response, or to stimulate or amplify an existing immune response. The term immunogenicity includes, for instance, allergenicity.

Within the context of the present invention, the term "allergenicity" designates the ability of a composition or protein or molecule to induce allergy in a mammal. This includes for instance the ability to trigger allergy, or to stimulate or amplify allergy. In a particular instance, allergenicity designates the ability to induce or stimulate production of IgE *in vivo*.

"Reducing" allergenicity or immunogenicity indicates a diminution in the allergenicity or immunogenicity, preferably by at least 20%, 30%, 40%, 50% or more. In preferred embodiment, "reducing" allergenicity or immunogenicity designates a reduction by at least 70%, 80%, 90% or more of the allergenicity or immunogenicity of a composition. The term "reducing" also encompasses the suppression of the allergenicity or immunogenicity.

"Removing" a component means at least reducing the amount of said component, preferably by at least 20% as compared to a reference material. In a particular embodiment of the invention, "removing" a component designates a removing of at least 50%, 60%, 70% or more of said component, even more preferably removing of at least 80%, at least 90%, at least 95%, 96%, 97%, 98%, 99%, or more. In a particular embodiment, removing encompasses com-

plete removal of a component, wherein the resulting material is substantially free of said component.

A "protein" designates a molecule comprising amino acids. The term thus designates polypeptides, proteins, or peptides, which may be of natural origin, purified, modified, recombinant, synthetic, etc. Peptides according to this invention typically contain between 3 to 70 amino acids in length, particularly from 5 to 50, from 5 to 40, or from 5 to 30.

A "cationic" protein means, within the context of the present invention, a protein having an isoelectric point of 7.4 or above, preferably of 7.6 or above, 7.8 or above, 8 or above, 8.5 or above, more preferably of 9 or above. A "cationic fraction" of a composition designates a fraction of that composition which contains cationic protein(s).

The term "allergen" designates any molecule that can cause allergy in a mammal. A proteinaceous allergen designates an allergen comprising amino acids in its structure.

The allergen according to the invention may be selected from food, respiratory, contact, or environmental allergens, for example, peanut, egg, milk or mite allergens. Examples of peanut allergens are selected among Ara h 1, Ara h 2, Ara h 3, Ara h 5, Ara h 6, Ara h 7, Ara h 8, Ara h 9, Ara h 10 and Ara h 11 peanut proteins.

The term "transcription infidelity" (TI) refers to a controlled mechanism by which DNA transcription generates RNA molecules with incorrect sequence(s), that subsequently leads to aberrant proteins. Transcription infidelity has been discovered and described by applicant in e.g., WO2008/009751. TI can generate RNAs with one or more differences. Such differences may be e.g., nucleotide substitutions, insertions and/or gaps (deletions), which eventually generate TI proteins with aberrant sequences. TI proteins of particular interest in the context of the present invention are TI gap proteins, which comprise a sequence resulting from suppression of a nucleotide during transcription, leading to proteins with a modified C-terminal end.

As described in example 1 of the present application, the applicant surprisingly observed that a protein normally incapable of inducing an immune response (IgE production) acquired this capability when it underwent a gap modification resulting from transcription infidelity. Thus, the presence of a C-terminal sequence resulting from the gap introduces into this protein a motif giving it an immunogenic (allergenic) nature absent from the normal protein.

To confirm the importance of this mechanism, the applicant carried out a bioinformatics annotation of all transcription infidelity events on the major milk proteins. In order to define precisely the appearance of a TI event, we thus carried out a next-generation sequencing (RNA-Seq) of the RNAs that encode milk proteins. Remarkably, the sequences obtained made it possible to identify exactly the positions of TI in milk proteins, and to know precisely the sequence of TI gap proteins (see example 2). In particular, we identified the following peptides, resulting from TI gaps, and conferring on the corresponding proteins an allergenic nature (Table 1):

Protein	Peptide Sequence	SEQ ID NO:
CSN1S1	LWHLFQKCLERRRSMN	1
CSN1S2	LPAFWLLPLQRIRWNMSPVVRNLLSSPRKHISR KRIWPLIPARRTFAPHSARKL	2

-continued

Protein	Peptide Sequence	SEQ ID NO:
CSN2	EAFQAVRNLLHASIRKLRSPFRVNRSSKQRMNS RIKSTPLPRHSL	3
CSN3	FWVPRSKTKNNQYAVRKMKDSSVTK	4
PAEP	KSTCSSAWRTVLSPSKAWPASAWSGPRRWTR PWRNSTKPSRPCPCTSGCPSTQPSWRSSATS	5

Continuing our research, we also surprisingly observed that TI proteins resulting from a shift in the reading frame by omission of a base in the RNA sequence (gap) have a higher content in positively-charged basic amino acids and a much lower content in acidic amino acids. This results in proteins having a cationic nature. As an illustration, the isoelectric point of the TI gap proteins identified in milk is given in the table 2 below:

	Protein			
	Canonical protein MW (Da)	Canonical protein pI	TI protein MW (Da)	TI protein pI
CSN1S1	22791	4.5	6418	11.2
CSN1S2	26019	8.6	7032	13.4
CSN2	29221	6.6	12196	11.6
CSN3	21269	6.7	4906	11.4
PAEP	19883	4.7	19684	9.3

From this table 2 it can be seen that TI gap proteins having an allergenic nature also have an isoelectric point much higher than that of the canonical proteins. This cationic nature of these proteins is advantageous because it makes it possible in particular to remove them by separation techniques based on isoelectric point. Accordingly, by cation-exchange technique, it is possible to remove proteins having an isoelectric point above a predefined value. Hence, it is possible to produce hypoallergenic (or low immunogenic) compositions by removing the cationic fraction therefrom. In this respect, as confirmed in example 3, milk depleted of the cationic fraction does not induce IgE in vivo, whereas the cationic fraction leads to massive IgE production.

The invention thus makes it possible to design new tools and processes for i) the early and precise molecular diagnosis of all forms of clinically significant allergies, ii) producing compositions having a less immunogenic/allergenic nature by removal of cationic proteins or TI-produced proteins, iii) removing TI-produced cationic proteins from certain food preparations in order to prevent or reduce the onset of allergies, iv) removing the main sources of allergens of certain food preparations intended for subjects identified as at risk, v) producing new adjuvants used to induce antibodies, or vi) defining new desensitization strategies capable of addressing all forms of allergies independently of their specificity.

An aspect of the invention thus relates to methods for reducing allergenicity or immunogenicity of a product by removing TI proteins therefrom, or by removing a cationic fraction therefrom. This method may be applied to various products such as food products (e.g., milk, peanut), cosmetic products, feed, pharmaceutical products, etc.

In this regard, an object of the invention relates to a method for preparing a food product comprising (i) providing a food product preparation, (ii) treating the food product

preparation to remove TI proteins therefrom, preferably TI gap proteins, and (iii) optionally formulating the treated food product with one or more suitable excipients.

Another object of the invention relates to a method for preparing a food product comprising (i) providing a food product preparation, (ii) treating the food product preparation to remove cationic proteins therefrom and (iii) optionally formulating the treated food product with one or more suitable excipients.

Another object of the invention relates to a method for preparing a cosmetic product comprising (i) providing a cosmetic agent, (ii) treating the cosmetic agent to remove TI proteins (preferably TI gap proteins) or cationic proteins therefrom and (iii) optionally formulating the treated cosmetic agent with one or more suitable excipients.

Another object of the invention relates to a method for preparing a pharmaceutical agent comprising (i) providing a pharmaceutical agent, (ii) treating the pharmaceutical agent to remove TI proteins (preferably TI gap proteins) or cationic proteins therefrom and (iii) optionally formulating the treated pharmaceutical agent with one or more suitable excipients.

In a preferred embodiment, the above methods comprise removing, from the products or compositions, cationic proteins that result from transcription infidelity, even more particularly from a transcription infidelity gap. More preferably, the method comprises removing at least 50% of cationic proteins having an isoelectric point of 7.4 or more.

In a preferred embodiment, the treated composition contains less than 2% by weight of cationic proteins having an isoelectric point above 8, more preferably less than 1%, even more preferably less than 0.5%, less than 0.2%, or less than 0.1%.

For instance, our analyses show that untreated milk contains approximately 3% by weight of cationic proteins having an isoelectric point above 7.4. The method of the invention preferably removes at least 50% of said proteins, thus leading to a milk preparation containing less than 1.5% by weight of cationic proteins. In a preferred embodiment, the treated milk preparation of the invention contains less than 1%, more preferably less than 0.5%, less than 0.4%, less than 0.3%, less than 0.2%, or even less than 0.1% by weight of cationic proteins having an isoelectric point above 8. The milk may be from any non-human mammal, such as cow, goat or sheep. The milk may also be artificial milk.

In this regard, the invention also relates to a food composition comprising a food product and a suitable excipient, wherein the food product contains less than 1% by weight of cationic proteins having an isoelectric point above 7.4, more preferably less than 0.5%, less than 0.3%, less than 0.2%, or less than 0.1%. In a most particular embodiment, the food product is milk, a cereal or peanut.

The invention also relates to a food composition comprising a food product and a suitable excipient, wherein the food product contains less than 1% by weight of proteins having a sequence resulting from TI gap, more preferably less than 0.5%, less than 0.3%, less than 0.2%, or less than 0.1%. In a most particular embodiment, the food product is milk, a cereal or peanut.

In a particular embodiment, the invention relates to a method for preparing a milk or a dairy product, comprising removing at least one protein comprising a TI gap peptide sequence selected from SEQ ID NO: 1 to 5.

In another particular embodiment, the invention relates to a method for preparing a peanut product, comprising removing at least one protein comprising a TI gap peptide sequence selected from SEQ ID NO: 110 to 120.

In a particular embodiment, the invention relates to a method for preparing a milk or a dairy product, comprising removing at least one protein comprising a sequence selected from SEQ ID NO: 6 to 10.

In another particular embodiment, the invention relates to a method for preparing a peanut product, comprising removing at least one protein comprising a sequence selected from SEQ ID NO: 121 to 131.

The invention also relates to a milk or a dairy product, or a peanut product, comprising less than 0.5% by weight of a protein comprising a TI gap peptide sequence selected from SEQ ID NO: 1 to 5 or SEQ ID NO: 110 to 120, respectively, preferably less than 0.3%, less than 0.2%, or less than 0.1%.

The invention also relates to a milk or a dairy product, or a peanut product, comprising less than 0.5% by weight of a protein comprising a sequence selected from SEQ ID NO: 6 to 10 or SEQ ID NO: 121 to 131, respectively, preferably less than 0.3%, less than 0.2%, or less than 0.1%.

The invention also relates to a cosmetic composition comprising a cosmetic agent and a suitable excipient, wherein the cosmetic agent contains less than 1% by weight of cationic proteins having an isoelectric point above 7.5, more preferably less than 0.5%, less than 0.3%, less than 0.2%, or less than 0.1%.

The invention also relates to a pharmaceutical composition comprising a pharmaceutical agent and a suitable excipient, wherein the pharmaceutical agent contains less than 1% by weight of cationic proteins having an isoelectric point above 7.5, more preferably less than 0.5%, less than 0.3%, less than 0.2%, or less than 0.1%.

The invention may also be used to produce immunogenic preparations having reduced allergenicity, suitable for desensitization of allergic subjects. In this regard, an object of the invention also relates to a method for preparing an allergen composition comprising (i) providing a proteinaceous allergen preparation, (ii) treating the preparation to remove cationic proteins therefrom and (iii) optionally formulating the preparation with one or more suitable excipients. Such treated preparations retain immunogenicity and may be used to induce tolerance in allergic subjects. However, such treated preparations have by themselves a reduced allergenicity and are less likely to induce undesirable side effects. Such method may be used with any allergen preparation, such as respiratory, contact, food, or environmental allergens (i.e., peanut, egg, milk, mite allergens, etc.). The allergen may be a recombinant proteinaceous allergen or a partially purified natural allergen.

The invention thus also relates to a composition comprising a proteinaceous allergen and a suitable excipient, wherein the proteinaceous allergen contains less than 2% by weight of cationic proteins, more preferably less than 1%, less than 0.5%, less than 0.2%, or less than 0.1%.

The invention also concerns a method of treating a subject allergic to an allergen to desensitize said subject to said allergen, comprising administering to the subject an effective amount of a composition as defined above.

The invention may also be used to produce pharmaceutical preparations having reduced allergenicity or immunogenicity. In this regard, an object of the invention also relates to a method for preparing a pharmaceutical composition comprising (i) providing a proteinaceous drug preparation, (ii) treating the preparation to remove cationic proteins therefrom and (iii) optionally formulating the preparation with one or more suitable excipients. Such treated preparations retain pharmaceutical activity and may be used to treat subjects. However, such treated preparations have by themselves a reduced allergenicity/immunogenicity and are less

likely to induce undesirable side effects. Such method may be used with any proteinaceous drug preparation, such as a drug, hormone, cytokine, enzyme, growth factor, etc.

As previously mentioned, removal of a cationic fraction or protein may be accomplished by techniques known per se in the art. In particular, cationic proteins may be removed by separation using cation exchange techniques. Examples of suitable material for cationic exchange include, for instance, HiTrap SP or CM, HiLoad SP or CM, or Bulk SP or CM. In a particular method, the process comprises (i) adjusting pH to a desired value (typically between 7.4 and 9) and (ii) subjecting the preparation to cationic exchange wherein essentially all material having an isoelectric point above the adjusted pH value is removed. In a particular embodiment, the method thus comprises (i) adjusting a solution of the composition to have a pH comprised between 7.4 and 9, preferably between 7.4 and 8.5, (ii) subjecting the solution to cation exchange chromatography, and (iii) recovering the eluate.

Alternatively, or in addition to the cationic exchange, the method may comprise a step of affinity chromatography using e.g., antibodies directed against transcription infidelity proteins. Such antibodies may be produced by procedures generally known in the art. For example, polyclonal antibodies may be produced by injecting TI proteins or a TI peptide thereof, or a cationic fraction of a biological sample, alone or coupled to a suitable carrier or adjuvant into a non-human animal. After an appropriate period, the animal is bled, sera recovered and purified by techniques known in the art (Paul, W. E. "Fundamental Immunology" Second Ed. Raven Press, NY, p. 176, 1989; Harlow et al. "Antibodies: A Laboratory Manual", CSH Press, 1988; Ward et al (Nature 341 (1989) 544).

The present invention also relates to a method for reducing immunogenicity or allergenicity of a peanut, the method comprising reducing the transcription infidelity rate in one or more peanut proteins selected from Ara h 1, Ara h 2, Ara h 3, Ara h 5, Ara h 7, Ara h 8, Ara h 9, Ara h 10 and Ara h 11. Such a method may comprise a step of integrating a synthetic DNA nucleotide sequence in a peanut grain, seed, or plant, wherein said synthetic sequence limits the transcription infidelity (TI) rate which results in TI gaps. Such a method may alternatively comprise a step of correcting and/or genetically modifying a DNA sequence in order to limit transcription infidelity rate and to avoid generation of TI peanut protein variants, by using various known gene engineering techniques such as Clustered Regularly Interspaced Short Palindromic Repeats (CRISPR) technology. In a particular embodiment, the invention relates to a transgenic peanut plant or a seed or grain thereof, comprising a modified Ara h protein gene with reduced transcription infidelity rate. The modified gene preferably comprises a modified T or A repeat domain.

The invention also allows the development of methods for detecting allergy in a subject. In this respect, the invention thus also concerns a method for detecting a subject having predisposition to allergy, comprising measuring in a sample from said subject the level of IgE directed against proteins having a sequence resulting from a transcription infidelity, wherein a difference in said level as compared to a control value indicates a subject having predisposition to allergy.

The invention also allows the stimulation of an immune response. Indeed, the identified TI proteins or peptides show increased immunogenicity and may be used to induce or stimulate an immune response, for instance as adjuvants.

The invention thus also relates to a cationic protein or peptide comprising a sequence resulting from transcription

infidelity, for use as an adjuvant to stimulate an immune response in a mammal, particularly to stimulate antibody production. The invention is particularly suited to stimulate or induce IgE production in a mammal. The mammal may be a non-human or a human mammal.

The invention also concerns a method of inducing or stimulating antibody production in a mammal, such as a human, comprising administering to the mammal a cationic protein or peptide having a sequence resulting from transcription infidelity.

The invention also concerns a vaccine composition comprising an immunogen and a protein or peptide having a sequence resulting from transcription infidelity.

A further object of the invention relates to a method for producing antibodies comprising (i) administering to a non-human mammal a cationic protein or peptide resulting from transcription infidelity, (ii) collecting antibodies produced and (iii) optionally deriving monoclonal and/or humanized antibodies from said collected antibodies.

It is also an object of the invention to provide methods of making anti-IgE antibodies, comprising (i) administering to a non-human mammal a cationic protein or peptide resulting from transcription infidelity under conditions allowing induction of antibody production, (ii) collecting antibodies produced and (iii) selecting antibodies that bind Fc receptor.

The invention also relates to a protein or peptide comprising a sequence selected from SEQ ID NO: 1-5 or SEQ ID NO: 74 to 109 or SEQ ID NO: 110 to 120, a fragment thereof, and to a composition comprising such a protein or peptide. The peptides, or fragments thereof, of the invention preferably have a length below 70 amino acid residues, even more preferably below 60, below 50, 40, below 35, or below 30. Particular peptides of the invention consist of anyone of SEQ ID NO: 1-5 or a fragment thereof containing at least 10 consecutive residues thereof. The fragments preferably shall retain an immunogenicity or allergenicity. The invention also relates to a protein comprising any one of SEQ ID NOs: 6-10 or SEQ ID NO: 16 to 51 or SEQ ID NO: 121 to 131.

Further aspects and advantages of the invention will be disclosed in the following experimental section, which is illustrative of the claims.

EXAMPLES

Example 1: Proof of Concept of the Molecular Origin of Allergy

The applicant has an approved animal facility for experimentation on mice and developed and published two pre-clinical murine models of allergy, one with peanut (Proust et al., 2008, *Int Arch Allergy Immunol* 146, 212-218), the other with cow's milk (Proust et al., 2009, *European Annals of Allergy and Clinical Immunology* 41(3): 85-94).

The hypothesis of the molecular origin of allergy was tested in the milk allergy model in which 6 successive intragastric administrations of proteins (at a rate of one per week) induce IgE production if the proteins are allergenic. Two different peptides were selected to test the hypothesis: a normal peptide not having undergone a TI event (non-TI peptide), and a TI peptide (resulting from a transcription infidelity event). These two peptides were administered to mice for 6 weeks and IgE production was followed over time (FIG. 1).

As shown in FIG. 1, IgE synthesis is massive when the mice are exposed to the TI peptide, whereas it is negligible

with non-TI peptide. Using a single intraperitoneal administration of TI and non-TI peptides, the same results were obtained (FIG. 2).

These two experiments show that TI peptides, by their specific physicochemical properties, are at the origin of IgE production, in both intragastric and intraperitoneal administration. This example illustrates that the large protein heterogeneity generated by transcription infidelity is a source of allergy triggers.

Example 2: Analysis of Milk Proteins

In order to obtain RNAs that encode milk proteins, we used the epithelial cells present in milk. Indeed, these cells are a good alternative for studying expression of RNAs expressed in bovine mammary gland (Canovas et al., 2014, *Scientific Reports* 4:5297). We obtained from a farm the milk of an untreated Prim'Holstein cow. Epithelial cells were purified from this milk.

From these cells, total RNA was extracted. The integrity of these RNAs was confirmed (RNA Integrity Number (RIN)=7.6) before preparing the libraries for Illumina next-generation sequencing. This RIN value is compatible with the preparation of Illumina libraries.

We have been able to confirm that the mean quality index for the two libraries is above 30 (library 1: 35.6 and library 2: 35.6) and that for the two libraries 94% of the reads have a quality index over 30. The quality of the sequences is compatible with the standards required for our analyses.

In order to define the positions where TI occurs, the reads obtained were aligned against the *Bos taurus* reference genome version UMD_3.1.1 (see Worldwide Website: ncbi.nlm.nih.gov/genome/82) and against the transcriptome constructed from the genome.

First, we studied the type of TI difference that leads to the most important modifications of the sequence of the protein encoded by the RNA. This is a deletion, also termed "TI gap" (FIG. 3).

When a TI gap appears in an RNA sequence, the protein sequence is highly modified. Indeed, the deletion of a base causes a shift of the reading frame, itself having dramatic consequences on the protein sequence located downstream of the event. Remarkably, we discovered that the carboxy-terminal part of the protein resulting from transcription infidelity (deletion) is enriched in basic amino acids and impoverished in acidic amino acids. Hence, the proteins resulting from TI have a cationic nature.

As proof of concept, we analyzed the sequences of the major milk allergens identified in the Allergome database (see Worldwide Website: allergome.org/). The allergens are presented in Table 3 below.

TABLE 3

List of major allergens in cow's milk.		
Gene	description	chromosome
ALB	albumin	6
CSN1S1	casein alpha s1	6
CSN1S2	casein alpha-S2	6
CSN2	casein beta	6
CSN3	casein kappa	6
LALBA	lactalbumin, alpha-	5
LPO	lactoperoxidase	19
LTF	lactotransferrin	22
PAEP	beta lactoglobuline	11

13

With the sequence data, we can estimate the expression of these genes in cells isolated from milk. To that end, we calculated the number of reads aligned to each gene (RPKM: Reads Per Kilobase per Million mapped reads) (Mortazavi et al., 2008, Nature Methods, 5(7): 621-28).

The results are shown in Table 4 below:

TABLE 4

Expression of genes encoding major allergens in cells isolated from cow's milk. Expression is measured in the 2 libraries in RPKM.

Gene	RPKM <i>moyen</i>	
	Librairie 1	Librairie 2
ALB	0.02	0.01
CSN1S1	131.91	128.72
CSN1S2	71.49	70.44
CSN2	153.00	154.07
CSN3	155.89	159.23
LALBA	39.12	43.29
LPO	0.13	0.11
LTF	10.63	10.91
PAEP	116.38	122.29

We selected the TI gap events that occur with high frequencies at the same positions in the two libraries. Similarly, the TI gaps that encode proteins having sequences the most enriched in basic amino acids were selected. We were thus able to identify the following TI proteins in milk (SEQ ID NOs: 6-10): the underlined part corresponds to the sequence of the TI gap peptide. For comparison, the sequences of the canonical proteins are provided (SEQ ID NOs: 11-15):

CSN1S1 (SEQ ID NO: 11)
 MKLLILTCLVAVALARPKHPIKHQGLPQEVLNENLLRFFVAFPFVEFGKE
 KVNELSKDIGSESTEDQAMEDIKQMEAESISSSEIIVPNSVEQKHQKED
 VPSERYLGYLEIVPNSAEERLHSMKEGIIHAQQKEPMIGVNQELAYFYPEL
 FRQFYQLDAYPSGAWYVPLGTQYTDAPSFSDIPNPIGSENSEKTTMPLW

CSN1S1_TI (SEQ ID NO: 6)
 MKLLILTCLVAVALARPKHPIKHQGLPQEVLNENLLRFLWHLWFQKCLERR
RSMN

CSN1S2 (SEQ ID NO: 12)
 MKFFIFTCLLAVALAKNTMEHVSSEESIISQETYKQEKMAINPSKENL
 CSTFCKEVRNANEEYSIGSSSESAEVATEEVKIVTDDKHQKALNEI
 NQFYQKFPQYLQYLYQGPIVLPNDQVQRNAVPIPTPLNREQLSTSEENS
 KKTVDMESTEVFTKTKLTKLEEKRNRLNPLKKISQRYQKFPALPQYLKTVYQ
 HQKAMKPWIQPKTKVIPYVRYL

CSN1S2_TI (SEQ ID NO: 7)
 MKFFILPAFWLLPLQIRIRWNMSPVVRNLSSPRKHISRKRIWFLIPARRTF
APHSARKL

CSN2 (SEQ ID NO: 13)
 MPLNTIYKQPQNQIIHSAPPSLLVLYPGKKELRAMKVLILACLVALALA
 RELEELNVPGEIVESLSSEESITRINKKIEKFQSEEQQTEDELQDKIH

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-continued

PFAQTQSLVYPPFGPIHNSLPQNIPLTQTPTVVVPPFLQPEVMGVSKVKE
 AMAPKHKEMPPFKYPVEPFTERQSLTLTDVENLHLPLPLLOSWMHQPHQ
 5 LPPTVMFPQSVLSLSQSKVLPVQKAVPYQORDMPIQAFLLYQEPVLRG
 VRGPFPIIV

CSN2_TI (SEQ ID NO: 8)
 10 MPLNTIYKQPQNQIIHSAPPSLLVLYPGKKELRAMKVLILACLVALALA
 RELEELNVPGEIVEAFQAVRNLLHASIRKLSRFRVRNSSKQRMNSRIKST
PLPRHSL

CSN3 (SEQ ID NO: 14)
 15 MMKSFLLVVTILALTLPLFGAQEQNQEQPIRCEKDERFFSDKIAKYIPIQ
 YVLSRYPYGLNYYQKQVALINNQLFLPYPIYAKPAAVRSPAQILQWQVL

20 SNTVPAKSCQAQPTTMRHPHPLSFMAIPPKKNQDKTEIPTINTIASGE
 PTSTPTTEAVESTVATLEDSPEVIESPEINTVQVTSTAV

CSN3_TI (SEQ ID NO: 9)
 25 MMKSFLLVVTILALTLPLFWVPRSKTKNNQYAVRKMKDSSTK
 PAEP (SEQ ID NO: 15)
 MKCLLLALALTCGAQALIVTQTMKGLDIQKVAGTWYSLAMAASDISLLDA
 QSAPLRVYVEELKPTPEGDLEILLQKWENGECAQKKIIAETKIPAVFKI
 30 DALNENKVLVLDTDYKYLFCMENSAPPEQSLACQCLVRTPEVDDEALE
 KFDKALKALPMHIRLSFNPTQLEEQCHI

PAEP_TI (SEQ ID NO: 10)
 35 MKCLLLALALTCGAQALIVTQTMKGLDIQKVAGTWYSLAMAASDISLLDA
 QSAPLRVYVEELKPTPEGDLEILLQKWENGECAQKKIIAETKIPAVFKI
 DALNENKVLVLDTDYKSTCSSAWRTVLSPSKAWPAWASGPRRWTRPWR
 40 NSTKPSRPCCTSGCPSTQPSWRSSATS

We analyzed the whole genome and confirmed that the disclosed peptides cannot be produced from another sequence in the cow transcriptome or genome.

45 The peptides thus identified, and the full-length proteins that contain them, are milk allergens. Removing these proteins from milk makes it possible to obtain a hypoallergenic milk. Furthermore, these proteins and peptides can also be used as adjuvants to stimulate an immune response in mammals, in particular to stimulate antibody production.

50 Example 3: Production of a Hypoallergenic Milk

The physicochemical properties of TI peptides and their importance in IgE production led us to develop a hypoallergenic infant formula that lacks these proteins.

55 For this purpose, we carried out a chromatographic fractionation of milk so as to produce two fractions: one enriched in TI proteins and the other depleted of TI proteins.

60 The initial rough material is in solid phase (whey powder). The material is composed of all soluble native milk proteins except caseins.

The material is dissolved in a buffered solution adjusted to the desired value (i.e. pH 7.4). The remaining insoluble parts are then removed by filtration or by centrifugation. The material is injected on a cationic exchange chromatography

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column (HiTrap SP FF) using the Akta Xpress chromatographic system (GE Healthcare Lifesciences). The separation is based on the isoelectric point of each protein. The proteins whose isoelectric points are higher than pH 7.4 (i.e. the cationic fraction) will bind to the resin, and the proteins whose isoelectric points are equal or lower than pH 7.4 (i.e. the non-charged and anionic fraction) will elute in the flow-through (FT). Once the FT is collected, the column is washed. The cationic fraction (which contains proteins with isoelectric points higher than pH 7.4) is then eluted from the column and collected by injecting a buffered solution of pH 7.4.

Therefore, the FT corresponds to the initial material depleted from its cationic fraction and is called Product no 1. Product no 1's main property is that 95% of cationic fraction which triggers allergy have been removed. Product no 1 is thus obtainable by a one-step physicochemical separation and is industrially compatible.

If suitable, a further treatment step may be performed to remove essentially all TI proteins. Such further treatment is an affinity binding step that uses Transcription Infidelity Antibodies (TIAB). More particularly, the TIABs are obtained from bovine sera. Briefly, the bovine serum is thermo-activated and then total IgGs are purified by affinity chromatography (based on Protein G). Product no 1 is incubated with the bovine TIABs and then submitted to an affinity chromatography based on Protein G. The total IgGs, including the ones complexed with proteins originating from Transcription Infidelity, bind to the resin and the FT contains a material totally depleted of allergy triggers (called Product no 2).

The fractions produced were administered via intragastric route to mice according to the protocol shown in FIG. 1 and IgE production was followed over time (FIG. 4).

The results presented in FIG. 4 show that mice exposed to the fraction enriched in TI proteins produce IgE in a massive way, whereas mice exposed to the depleted fraction produce little or no IgE. We can thereby conclude that 1) TI proteins in milk are indeed at the origin of milk's allergenicity and 2) the depleted fraction is much less allergenic than the enriched fraction. The invention thus makes it possible to produce a hypoallergenic milk.

Example 4: Analysis of Mite Allergens

This analysis is based on the localization of TI gap events in mite allergens and non-allergens. We have surprisingly observed that TI deletion events are mainly present in the coding regions (ORF-IN) of the transcripts coding for mite allergens whereas TI deletion events are mainly outside the coding regions (ORF-OUT) in the case of transcripts which do not code for mite allergens.

Der f 1_iso1

MKFVLAIASLLVLS TVYARPASIKTFEEFKKAFNKNYATVEEEVARKNFLES L
 KYVEANKGAINHLS DLSDLEDFKNRYLMSAEAFQ LKTQFDLNAETSACRINSVN
 VPSELDLRLSLRTVTPIRMQGGCGSCWAFSGVAATESAYLAYRNTSLDLSEQELV
 DCASQHGCHGDTIPRGI EYIQQNGVVEERSYPYVAREQQCRRPNSQHYGISNYC
 QIYPPDVKQIREAL TQHTAI AVI IIGIKDLRAFQHYDGR TIIQHDNGYQPNYHAV
 NIVGYGSTQGVVDYIWRNSWDTTWGDSGYGYFQAGNNLMMIEQYPYVIVM

Der f 1_iso1_TI

MKFVLAIASLLVLS TVYARPASIKTFEEFKKAFNKNYATVEEEVARKNFLES L
 KYVEANKGAINHLS DLSDLEDFKNRYLMSAEAFQ LKTQFDLNAETSACRINSVN

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In order to carry out this study, we have analyzed RNA sequences of *Dermatophagoides farinae* by Illumina next-generation sequencing. We formed two groups of sequences: (1) Transcripts known as allergens and (2) Transcripts which are not allergens. The two groups have similar characteristics of sequence (such as frequency of four bases, mean coverage at each position, length of covered transcripts, exons number) as summarized in Table 5.

TABLE 5

Comparison of transcripts coding allergens versus transcripts coding non-allergens (i.e., proteins never described as allergens). Each group includes 35 transcripts. Evaluated parameters show that the two groups are similar.

	Allergen	Non allergen
Number of transcripts	35	35
A frequency	33.2 (±3.1)	34.1 (±2.6)
T frequency	29.8 (±2.5)	32.11 (±3.1)
C frequency	18 (±2.9)	17 (±2.1)
G frequency	18.9 (±2.3)	16.8 (±2.3)
Length of covered transcripts	666.5 (±446.3)	674 (±461.9)
Mean coverage at each position	61.8 (±62.1)	64 (±67.9)
Number of exons	3 (±1.6)	3 (±1.3)

We have identified the events of transcription infidelity (TI) in these transcripts and we have compared the two groups, namely the allergens and non-allergens. Our results show that TI deletions affecting the sequences of allergens are mainly located within the coding regions, i.e. in the ORF, contrary to TI deletion events affecting the sequences of non-allergens, which are mainly found in non-coding regions (see FIG. 5).

We have also analyzed a ratio between the number of deletions in the coding sequences and the number of deletions in the non-coding sequences, and we have observed that the deletions in the coding sequences of allergens preferentially affect repetitions of A or T base (see FIG. 6).

The repetitions of affected A bases are mainly followed by T bases and the repetitions of affected T bases are followed by G bases; these two types of events, when they are located in the ORF, are thus very specific to allergens. These events affect a considerable part of transcripts, from about 0.2% to 10%, more preferably from 0.2% to 5%.

We were able to identify the following TI allergenic proteins for all TI gaps events in *Dermatophagoides farinae*: SEQ ID NO: 16-51. The underlined parts of the sequences correspond to the sequences of the TI gap peptides also listed below as SEQ ID NO: 74 to 109. For comparison, the sequences of the canonical proteins are also provided (SEQ ID NO: 52-73).

(SEQ ID NO: 52)

(SEQ ID NO: 16)

-continued

VPSELDRSLRVTPTIRMQGGCGSCWAFSGVAATESAFWPTVTRLWIFLNRNSS
IAHLNTDVTAIQYQEASNTSNKMVSLKKEAIIHTLHENNNADDQIRNITVTSQTTA
KFIHQM

Der f 1_iso1_TI

(SEQ ID NO: 17)

MKFVLAIASLLVLS TVYARPASIKTFEEFKKAFNKNYATVEEEVARKNFLES
KYVEANKGAINHLS DLSLDEFKNRYLMSAEAFEQLKTOFDLNAETSACRINSVN
VPSELDRSLRVTPTIRMQGGCGSCWAFSGVAATESAYLAYRNTSLDLSSEQELV
DCASQHGCHGDTIPRGI EYIQQNGVVEERSYPVAREQQCRRPNSQHYGISNYC
QIYPPDVVKQIREAL TQHTAI AVIIGIKDLRASNIMMDEQSFNMTMVINQTIMPST
LSVTEVHKASII GSYETVGIQPGVIADTDISKPETTS

Der f 1_iso1_TI

(SEQ ID NO: 18)

MKFVLAIASLLVLS TVYARPASIKTFEEFKKAFNKNYATVEEEVARKNFLES
KYVEANKGAINHLS DLSLDEFKNRYLMSAEAFEQLKTOFDLNAETSACRINSVN
VPSELDRSLRVTPTIRMQGGCGSCWAFSGVAATESAYLAYRNTSLDLSSEQELV
DCASQHGCHGDTIPRGI EYIQQNGVVEERSYPVAREQQCRRPNSQHYGISNYC
QIYPPDVVKQIREAL TQHTAI AVIIGIKDLRAFQHYDGRITIIQHDMVINQTIMPSTL
SVTEVHKASII GSYETVGIQPGVIADTDISKPETTS

Der f 1_iso1_TI

(SEQ ID NO: 19)

MKFVLAIASLLVLS TVYARPASIKTFEEFKKAFNKNYATVEEEVARKNFLES
KYVEANKGAINHLS DLSLDEFKNRYLMSAEAFEQLKTOFDLNAETSACRINSVN
VPSELDRSLRVTPTIRMQGGCGSCWAFSGVAATESAYLAYRNTSLDLSSEQELV
DCASQHGCHGDTIPRGI EYIQQNGVVEERSYPVAREQQCRRPNSQHYGISNYC
QIYPPDVVKQIREAL TQHTAI AVIIGIKDLRAFQHYDGRITIIQHDNGYQPNYHAV
NIVGYGSTQGVDIGSYETVGIQPGVIADTDISKPETTS

Der f 1_iso2

(SEQ ID NO: 53)

MKFVLAIASLLVLS TVYARPASIKTFEEFKKAFNKNYATVEEEVARKNFLES
KYVEANKGAINHLS DLSLDEFKNRYLMSAEAFEQLKTOFDLNAETSACRINSVN
VPSELDRSLRVTPTIRMQGGCGSCWAFSGVAATESAYLAYRNTSLDLSSEQELV
DCASQHGCHGDTIPRGI EYIQQNGVVEERSYPVAREQQCRRPNSQHYGISNYC
QIYPPDVVKQIREAL TQHTAI AVIIGIKDLRAFQHYDGRITIIQHDNGYQPNYHAV
NIVGYGSTQGVYDVIWRNSWDTTWGDSGYGYFQAGNNLMMIEQYPPYVIM

Der f 1_iso2_TI

(SEQ ID NO: 20)

MKFVLAIASLLVLS TVYARPASIKTFEEFKKAFNKNYATVEEEVARKNFLES
KYVEANKGAINHLS DLSLDEFKNRYLMSAEAFEQLKTOFDLNAETSACRINSVN
VPSELDRSLRVTPTIRMQGGCGSCWAFSGVAATESAFWPTVTRLWIFLNRNSS
IAHLNTDVTAIQYQEASNTSNKMVSLKKEAIIHTLHENNNADDQIRNITVTSQTTA
KFIHQM

Der f 1_iso2_TI

(SEQ ID NO: 21)

MKFVLAIASLLVLS TVYARPASIKTFEEFKKAFNKNYATVEEEVARKNFLES
KYVEANKGAINHLS DLSLDEFKNRYLMSAEAFEQLKTOFDLNAETSACRINSVN
VPSELDRSLRVTPTIRMQGGCGSCWAFSGVAATESAYLAYRNTSLDLSSEQELV
DCASQHGCHGDTIPRGI EYIQQNGVVEERSYPVAREQQCRRPNSQHYGISNYC
QIYPPDVVKQIREAL TQHTAI AVIIGIKDLRASNIMMDEQSFNMTMVINQTIMPST
LSVTEVHKASII GSYETVGIQPGVIADTDISKPETTS

Der f 1_iso2_TI

(SEQ ID NO: 22)

MKFVLAIASLLVLS TVYARPASIKTFEEFKKAFNKNYATVEEEVARKNFLES
KYVEANKGAINHLS DLSLDEFKNRYLMSAEAFEQLKTOFDLNAETSACRINSVN
VPSELDRSLRVTPTIRMQGGCGSCWAFSGVAATESAYLAYRNTSLDLSSEQELV
DCASQHGCHGDTIPRGI EYIQQNGVVEERSYPVAREQQCRRPNSQHYGISNYC
QIYPPDVVKQIREAL TQHTAI AVIIGIKDLRAFQHYDGRITIIQHDMVINQTIMPSTL
SVTEVHKASII GSYETVGIQPGVIADTDISKPETTS

Der f 1_iso2_TI

(SEQ ID NO: 23)

MKFVLAIASLLVLS TVYARPASIKTFEEFKKAFNKNYATVEEEVARKNFLES
KYVEANKGAINHLS DLSLDEFKNRYLMSAEAFEQLKTOFDLNAETSACRINSVN
VPSELDRSLRVTPTIRMQGGCGSCWAFSGVAATESAYLAYRNTSLDLSSEQELV
DCASQHGCHGDTIPRGI EYIQQNGVVEERSYPVAREQQCRRPNSQHYGISNYC
QIYPPDVVKQIREAL TQHTAI AVIIGIKDLRAFQHYDGRITIIQHDNGYQPNYHAV
NIVGYGSTQGVDIGSYETVGIQPGVIADTDISKPETTS

Der f 23_iso1

(SEQ ID NO: 54)

MKNITIAFVSLAILIHSSYADIDHFDNDDQNSSSRPDDPTMIDVQTTTVQPS
SMPTTSESQSTVKPTTTTVKPSPTTVKLT TTTTVKPTTTTVKPTTTTVKPSPTTVKPT
TTTVKPSPTTT TTTTTPQPEDEFECPTRFYFADPKDCKFYICSNWEAIHKSCP
GNTRWNEKELTCT

-continued

Der f 23_isol_TI

(SEQ ID NO: 24)

MKFNTIAFVSLAILIHSSYADIDHFDNDDQNSSTRPDDPTTMIDVQTTTVQPS
SMPTTSESQSTVKPTTTTVKPSPTTVKLTTTTTVKPTTTTVKPTTTTVKPSPTTVK
TTTTVKPSPTTTTTTTEQPEDFEFCPTFRGFADPKDCKFIFVQIGKLYIKVVQ
VIQDGMKKN

Der f 11_isol

(SEQ ID NO: 55)

MSARTAKYMYRSSGAGASGDI SVEYGTDLGALTRLEDKIRLLSDDLESEEREMR
QRIEREKAELQIQVMSLGERLEEAEGSSSESVTEMNKKRDS ELAKLRKLLLEDVHI
ESEETAHHLRQKHQAAIQEMQDQLDQLQAKNKS DKEKQKQAEVFELLAQL
ETANKEKLTALKNVKLEYTVHELNIKIEEINRTVIELTSHKQRLSQENTEELI KEV
HEVKLQLDNANHLKTQIAQQL EDTRHRL EEEERKRASLENHAHTLEVELES LK
VQLDEESEARLELERQLTKANGDAASWKS KYEAELQAHADVEEELRRKMAQK
ISEYEEQLEALLNKCSSLEKQKSRLOSEVEV LMDLEKATAHAQQL EKRVQLE
KINLDDLKKNLEEV TMLMEQAQKELRVKIAELQKLQHEYEKLRDQRDQLAREN
KKLTD DLAEAKS QLNDAHRR IHEQEIEIKRLENERDEL SAAYKEAETLRKQEEA
KNQRLIAELAQRVHDY EKRLAQKDEEIEALRKQYQIEI EQLNMR LAEAEAKLKT
ETARLKKYQAQIT ELELSDAANKANIDLQKTIKKQALQITSELQAHYDEVHR
QLQQAVDQLGVTQRRCQALQAELEEMRIAL EQANRAKRQAEQLHEEA VVRVN
ELTTINVLNLSAKSKLESEFSALQADYDEVHKELRISDERVQKLTIELKSTKDLLI
EEQERLVKLETVKKSLQEVRTLHVRIEEVEANALAGGKRVIAKLESRI RDVETE
VEEERRRHAETDKMLRKKDHRVKELLLQNEEDHKQIQLLQEMTDKLNKVKV
YKRMQEQEGMSQQNLTRVRRFQRELEAAEDRADQAESNLSFIRAKHRSWVT
TSQVPGGTRQVFTTQEETNY

Der f 11_isol_TI

(SEQ ID NO: 25)

MSARTAKYMYRSSGAGASGDI SVEYGTDLGALTRLEDKIRLLSDDLESEEREMR
QRIEREKAELQIQVMSLGERLEEAEGSSSESVTEMNKKRDS ELAKLRKLLLEDVHI
ESEETAHHLRQKHQAAIQEMQDQLDQLQAKNKS DKEKQKQAEVFELLAQL
ETANKEKLTALKNVKLEYTVHELNIKIEEINRTVIELTSHKQRLSQENTEELI KEV
HEVKLQLDNANHLKTQIAQQL EDTRHRL EEEERKRASLENHAHTLEVELES LK
VQLDEESEARLELERQLTKANGDAASWKS KYEAELQAHADVEEELRRKMAQK
ISEYEEQLEALLNKCSSLEKQKSRLOSEVEV LMDLEKATAHAQQL EKRVQLE
KINLDDLKKNLEEV TMLMEQAQKELRVKIAELQKLQHEYENYVINV INWHVKT
NLQTLPKLNHN

Der f 11_isol_TI

(SEQ ID NO: 26)

MSARTAKYMYRSSGAGASGDI SVEYGTDLGALTRLEDKIRLLSDDLESEEREMR
QRIEREKAELQIQVMSLGERLEEAEGSSSESVTEMNKKRDS ELAKLRKLLLEDVHI
ESEETAHHLRQKHQAAIQEMQDQLDQLQAKNKS DKEKQKQAEVFELLAQL
ETANKEKLTALKNVKLEYTVHELNIKIEEINRTVIELTSHKQRLSQENTEELI KEV
HEVKLQLDNANHLKTQIAQQL EDTRHRL EEEERKRASLENHAHTLEVELES LK
VQLDEESEARLELERQLTKANGDAASWKS KYEAELQAHADVEEELRRKMAQK
ISEYEEQLEALLNKCSSLEKQKSRLOSEVEV LMDLEKATAHAQQL EKRVQLE
KINLDDLKKNLEEV TMLMEQAQKELRVKIAELQKLQHEYEKLRDQRDQLAREN
KKLTD DLAEAKS QLNDAHRR IHEQEIEIKRLENERDEL SAAYKEAETLRKQEEA
KNQRLIAELAQRVHDY EKRLAQKDEEIEALRKQYQIEI EQLNMR LAEAEAKLKT
ETARLKKYQAQIT ELELSDAANKANIDLQKTIKKQALQITSELQAHYDEVHR
QLQQAVDQLGVTQRRCQALQAELEEMRIAL EQANRAKRQAEQLHEEA VVRVN
ELTTINVLNLSAKSKLESEFSALQADTMKYIKNLEFLMNEYRNLQLN SNLLKIC

Der f 15_isol

(SEQ ID NO: 56)

MKTIYAILSIMACIGLMNASIKRDHNDYSKNPMRIVCYVGTWSVYHKVDPYTI E
DIDPFKCTHLMYGFAKIDEYKYTIQVFDPYQDDNHNSWEKRGYERFNNLR LKN
PELTMTISLGGWYEGSEKYS DMAANPTYRQQFIQSVLDFLQYKFDGLDLDWE
YPGSRLGNPKIDKQNYLALVRELKDAFEPHGYLLTAAVSPGKDKIDRAYDIKEL
NKLFDWMNVMTYDYHGGWENFYGHNAPLYKRPDETD ELHTYFNVNYTMHY
YLNNGATRDKLVMGVPPFYGRAWSIEDRSK LKLGDPAGKMSPPGFI SGBEGVLS
YIELCQLFPQKEEWHIQYDEYINAPYGYNDKIWVGDDLASISCKLAF LKELGV
SGVIVWLENDDFKGCHPKNPLLNKVNMMINGDEKNSFECILGPSTTTPTPTTT
PTTTPTSPPT
TTEHTSETPKYTT YVDGHLI KCYKEGDI PHPTN IHKYLVC EFVNGGWVHIMP
CPPGTIWCQEKLT CIGE

Der f 15_isol_TI

(SEQ ID NO: 27)

MKTIYAILSIMACIGLMNASIKRDHNDYSKNPMRIVCYVGTWSVYHKVDPYTI E
DIDPFKCTHLMYGFAKIDEYKYTIQVFDPYQDDNHNSWEKRGYERFNNLR LKN
PELTMTISLGGWYEGSEKYS DMAANPTYRQQFIQSVLDFLQYKFDGLD WIGSI
LDDLWVTRKSTNKT IWLWLENLKTLLNLMATC

Der f 32_isol

(SEQ ID NO: 57)

MSTTNYSDVHRGFSNLDYRIYFKDNSNGKI I SPWHDIP LFVDKSAKHYNMVVE
IPRWTEKMEIATAEPMSPIKQDIKKGALRVVKNVFP HKGYIWNYGAPQTWE

-continued

NPNHIDQDTKTKGDNNDPIDVIEIGSRVAKRGDVPVKILGTIALIDEGETDWKIIA
IDTRDELASQMNNDVDEKLLPGLLRATVWEFKIYKIPDGKPAKFAFNGEAK
DREFAEKIVEETHQYWQEMMENKSGEHKLDLKNVTLGNSFSINDEQAKQFLET
RPSSDAVEPTPIADQVAIDKWHEIVKLI

Der f 32_isol_TI

(SEQ ID NO: 28)

MSTTNSVDHRGFSNLDYRIYFKDNSNGKII SPWHDIPLFVDKSAKHYNMVE
IPRWITNEKMEIATAEPMSPFKQDIKKGALRYVKNVFPKGYIWNYGAPPQTWE
NPNHIDQDTKTKGDNNDPIDVIEIGSRVAKRGDVPVKILGTIALIDEGETDWKIIA
IDTRDELASQMNNDVDEKLLPGLLRATVWEFKIYKIPDGKPAKFAFNGEAK
DRELLKKS LKKHINIGKK

Der f 25_isol

(SEQ ID NO: 58)

MVRKFFVGGNWKMNNGSRATNEDLIKTLNGLDPLDNTDVPVGVPSIYMAEVRQ
KLPKTI GVAQAQNCYKVPKGAPTGEI SPAMI KDVGAEWVILGHSERRNVFGESDQ
LIGEKVEHALQEGHLVIAACIGELLEEREAGKTTEVVFRTQVISKHVKDWSKVV
LAYEPVWAIGTGKTASPQQAQEVHQKLRQWFSENVSPQIAETIRIYGGSVTAN
NAKELASQADVDDGFLVGGASLKEPEFVQVFNARQ

Der f 25_isol_TI

(SEQ ID NO: 29)

MVRKFFVGGNWKMNNGSRATNEDLIKTLNGLDPLDNTDVPVGVPSIYMAEVRQ
KLPKTI GVAQAQNCYKVPKGAPTGEI SPAMI KDVGAEWVILGHSERRNVFGESDQ
LIGEKVEHALQEGHLVIAACIGELLEEREAGKTTEVVFRTQVISKHVKDWSKVV
LAYEPVGLVVLVKQPVHNKHKKFIKFNFDNGFLKMFHHLKPKQFESFMVVQ

Der f 25_isol_TI

(SEQ ID NO: 30)

MVRKFFVGGNWKMNNGSRATNEDLIKTLNGLDPLDNTDVPVGVPSIYMAEVRQ
KLPKTI GVAQAQNCYKVPKGAPTGEI SPAMI KDVGAEWVILGHSERRNVFGESDQ
LIGEKVEHALQEGHLVIAACIGELLEEREAGKTTEVVFRTQVISKHVKDWSKVV
LAYEPVWAIGTGKTASPQQAQEVHQKLRQWFSEMFHHLKPKQFESFMVVQ

Der f 16_isol

(SEQ ID NO: 59)

MAAHDKNFDVIPIGHTFFFIWRIKQFELVVPVKEDYKGFYKGDYIVACCTENP
TGGHSMESKPI LNHGYCHI HFWIGSESTKDEAGVAAIKSVELDDFLGGYPVQ
HREIEEFESRQFSSYFKNGI IYKGGYESGFTK MIDELKPSLLHVKGKRRPIVYEC
AEISWKVMNNGDVP ILLVNPVFWVTGKHSNRMERTTAIRVANDLKS ELNRFK
LSSVILEDGKEVEQTS GAEDAFNKALSLDKKIDIDLKQMPKGYDYAASDKSFE
SHERSFVTLKCFEGTETIDI SFVKNGPLSRADLDTNDTFIVENGSEGLWVWVG
KKATQKERQSAIKYAMELINKKYPNNTPVTKVLEGEDESVEFKSLFESWQMS
QEKI TSARLFRVSRNGIFKQVANYEPDDEEDNIMILDVMDKIYVWIGNQFAERI
ADEAHVDKVAQRFIQEDKSGRFRPNQIIKCLKQGSSEDAFKSYFPKWN

Der f 16_isol_TI

(SEQ ID NO: 31)

MAAHDKNFDVIPIGHTFFFIWRIKQFELVVPVKEDYKGFYKGDYIVACCTENP
TGGHSMESKPI LNHGYCHI HFWIGSESTKDEAGVAAIKSVELDDFLGGYPVQ
HREIEEFESRQFSSYFKNGI IYKGGYESGFTK MIDELKPSLLHVKGKRRPIVYEC
AEISWKVMNNGDVP ILLVNPVFWVTGKHSNRMERTTAIRVANDLKS ELNRFK
LSSVILEDGKEVEQTS GAEDAFNKALSLDKKIDIDLKQMPKGYDYAASDKSFE
SHERSFVTLKCFEGTETIDI SFVKNGPLSRADLDTNDTFIVENGSEGLWVWVG
KKATQKERQSAIKYAMELINKKYPNNTPVTKVLEGEDESVEFKSLFESWQMS
QEKI TSARLFRVSRNGIFKQVANYEPDDEEDNIMILDVMDKIYVWIGNQFAERI
ADEAHVDKVAQRFLYKRIKVAVNFDQIRL

Der f 26_isol

(SEQ ID NO: 60)

MALPRVFFDIAADNQPLGRIVIELRSDVVPKTAENFRALCTGEKGFQKSSSFHR
IIPNFMIOGGDFTNHNGTGGKSIYGNKFADENFLLQHTGPGIMSMANAGPNING
SQFFITTVKTTWLDGKHVVFGSVVEGMDIVKKVESYGSQSGKPSKKVTIANCG
QL

Der f 26_isol_TI

(SEQ ID NO: 32)

MALPRVFFDIAADNQPLGRIVIELRSDVVPKTAEISVHFALVKKDLVLNHPHFIV
SYPII

Der f 26_isol_TI

(SEQ ID NO: 33)

MALPRVFFDIAADNQPLGRIVIELRSDVVPKTAENFRALCTGEKGFQKSSSFHR
IIPNFMIOGGDFTNHNGTGGKSIYGNKFADENFLLQHTGPGIMSMANAGPNING
SQFFITTVKTTWLDGKHVVFGSVVEGMDIVKRWKMAHNRVNHPRK

Der f 23_isol2

(SEQ ID NO: 61)

MKFNITIAFVSLAILHSSYADIDHFDNDQNSSSRPDDPTMIDVQTTTVQPS
DEFECPTRFYFADPKDPCKFYICSNWEAIIHKSCPGNTRWNEKELTCT

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Der f 23_iso2_TI (SEQ ID NO: 34)
 MKFNITIAFVSLAILIHSSYADIDHFDNDDQNSSTSRPDDPTMIDVQTTTVPQS
 DEFECPTRFGYFADPKDPCKFIFVQIGKLYIKVVQVIQDGMKKN

Der f 13_iso1 (SEQ ID NO: 62)
 MASI EGKYKLEKSEKFEFLDKLGVGFMVKTAAKTLKPTFEVAIENDQYIFRSL
 STFKNTEAKFKLGEFEEDRADGKRKVTVIQKEGDNKVFQTFQGDKEVKIIREF
 NGDEVVVTASCDGVTSVRTYKRI

Der f 13_iso1_TI (SEQ ID NO: 35)
 MASI EGKYKLEKSEKFEFLDKLGVGFMVKTAAKTLKPTFEVAIENDQYIFRSL
 STFKNTEAKFKLGEFEEDRADGKRKVTVIQKVTINLFLKHNSVIKK

Der f 7_iso1 (SEQ ID NO: 63)
 MMKFLIIAAVAFVAVSADPIHYDKITTEEINKAIDDAIAAIEKSETIDPMKVPDHT
 DKFERHVGILDFKGLAMRNI EARGLKQMKRQGDANVKGEEGIVKAHLLIGVH
 DDIVSMEYDLAYKLGDLHPNTHVISDIQDFVVALSLEISDEGNITMTSFEVRQFA
 NVVNHIGGLSILDP IFGVLSVLTAFIQDQTVRKEMTKVLAPAFKRELEKN

Der f 7_iso1_TI (SEQ ID NO: 36)
 MMKFLIIAAVAFVAVSADPIHYDKITTEEINKAIDDAIAAIEKSETIDPMKVPDLPI
 NSNVMLVFWISKVN

Der f 27_iso1 (SEQ ID NO: 64)
 MKFLLSFVLMIVAATATYAAHVGSGSRDNNNNKVPVPAEGFAKASNEFGPHLL
 KEVIQHRSSSGSRGSENVLFSYVAVALSMVHQGTQGSTAEQFKRVLYYDR
 VQQLNGGEYQTVANSVKIQIQSDQSNQFDWGNMLMVDQOIPVKDQYK
 KIIIEQYYDQVMSVDFRKEKSNVMERINQFVSNKTHGLIDRMLQPPSADTGLA
 LINAVYFKGEWLKPFDSMRTEQSVFYGHGQYKQVYINGQGPYGYVEVPQ
 WNSDLIQLPYKGEDIAFYGVLPREPRNYDLDKIRQINSSTFVDEIVGQITGSQSSV
 YFPKIELSTSYQLPEILKSMGLQDVFTEADLSGITDKKPKMIDDIAHKAKLILNE
 QGTEAGAGTYIQMAVLSALETSHTRFRDHPMYFIRHLPTGGIILFLGEIHD

Der f 27_iso1_TI (SEQ ID NO: 37)
 MKFLLSFVLMIVAATATYAAHVGSGSRDNNNNKVPVPAEGFAKASNEFGPHLL
 KEVIQHRSSSGSRGSENVLFSYVAVALSMVHQGTQGSTAEQFKRVLYYDR
 VQQLNGGEYQTVANSVKIQIQSDQSNQFDWGNMLMVDQOIPVKDQYK
 KIIIEQYYDQVMSVDFRKEKSNVMERINQFVSNKTHGLIDRMLQPPSADTGLA
 LINAVYFKGEWLKPFDSMRTEQSVFYGHGQYKQVYINGQGPYGYVEVPQ
 WNSDLIQLPYKGEDIAFYGVLPREPRNMLTKFVNQSIQLLLMKLLDKLLVSHQ
 LFISRKLNSVHHINCRKF

Der f 7_iso2 (SEQ ID NO: 65)
 MMKFLIIAAVAFVAVSADPIHYDKITTEEINKAIDDAIAAIEKSETIDPMKVPDHT
 DKFERHVGILDFKGLAMRNI EARGLKQMKRQGDANVKGEEGIVKAHLLIGVH
 DDIVSMEYDLAYKLGDLHPNTHVISDIQDFVVALSLEISDEGNITMTSFEVRQFA
 NVVNHIGGLSILDP IFGVLSVLTAFIQDQTVRKEMTKVLAPAFKRELEKN

Der f 7_iso2_TI (SEQ ID NO: 38)
 MMKFLIIAAVAFVAVSADPIHYDKITTEEINKAIDDAIAAIEKSETIDPMKVPDLPI
 NSNVMLVFWISKVN

Der f 29_iso1 (SEQ ID NO: 66)
 MALPRVFFDIAADNQLGRIVIELRSDVVPKTAENFRALCTGEGKGFKSSSFHR
 IIPNFMIOGGDFTNHNGTGGKSIYGNKFADENFTLQHTGPGIMSMANAGPNTNG
 SQFFITTVKTTWLDGKHVVFGSVVEGMDIVKKVESYGSQSGKPSKKTIANCG
 QL

Der f 29_iso1_TI (SEQ ID NO: 39)
 MALPRVFFDIAADNQLGRIVIELRSDVVPKTAENFRALCTGEGKGFKSSSFHR
 IIPNFMIOGGDFTNHNGTGGKSIYGNKFADENFTLQHTGPGIMSMANAGPNTNG
 SQFFITTVKTTWLDGKHVVFGSVVEGMDIVKRWKAMAHNRVNHPRK

Der f 18_iso1 (SEQ ID NO: 67)
 MTRFSLTVLAVLAACFGSNIREPNVATLEPKTVCYYESVWHWRQEGKMDPEDI
 DTSLCTHIVSYFGIDAATHEIKLLDEYLMKDLHDMEHFTQHKGNAKAMIAVG
 GSTMSDQPSKTAAVEHYRETFFVSTVDLMTRYGFDGVMIDWSGMQAKDSNF
 IKLLDKFDEKFAHTSFVMGVTLPATIASYDNYNIPAINYVDFMNVLSLDYTG

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WAHTVGHASPFPEQLKTLEAYHKGAPRHKMVMVAPFYARTWILEKMNKQDI
GDKASGPGPRGQFTQTDGFLSYNELCVQIQAETNAFTI TRDHDNTAI YAVVHVS
NHAEWI SFEDRHTLGEKAKNI TQQGYAGMSVYTL SNEDVHGVC GDNPLLHAI
QSNYHYGVVTEPTVVTLPPVTHTEHVTDI PGVPHCHEEGFFRDKTYCATYYEC
KKGDFGLEKTVHHCANHLQAFDEVSRCTIDHTKIPGC

Der f 18_iso1_TI

(SEQ ID NO: 40)

MTRFSLTVLAVLAACFGSNIRPNVATLEPKTVCYYESVWHWRQGEKMDPEDI
DTSLCTHIVVSYFGIDAATHEIKLLEDEYLMKDLHDMEHFTQHKGNAKAMIAVG
GSTMSDQFSKTAAVEHYRETFVSTVDLMTRYGPDGVMIDWSGMQAKDSNDF
IKLLDKFDEKFAHTSFVMGVTLPATIASYDNYNI PAISNYVDFMNVLSLDYTG
WAHTVGHASPFPEQLKTLEAYHKGAPRHKMVMVAPFYARTWILEKMNKQDI
GDKASGPGPRGQFTQTDGFLSYNELCVQIQAETNAFTI TRDHDNTAI YAVVHVS
NHAEWI SFEDRHTLGEKAKNI TQQGYAGMSVYTL SNEDVHGVC GDNPLLHAI
QSNYHYGVVTEPTVVTLPPVTHTEHVTDI PGVPHCHEEGFFRDKTYCATYYEC
KKGDFGLEKPCIIIVPITYRHLLTK

Der f 16_iso2

(SEQ ID NO: 68)

MAAHDKNPDVIPIGHTFFFIWRIKQFELVVPKEDYGKFKGDCYIVACCTENP
TGGHSMKESKPI LN GHGYCHIFWIGSESTKDEAGVAAIKSVELDDFLGGYPVQ
HREIEEFESRQFSSYPKNGI IYKGGYESGPTKMIDELKPSLLHVKGKRRPIVYEC
AHSWKVMNNGDVFILLVFNPFVWVTGKHSNRMERTTAIRVANDLKS ELNRFK
LSSVILEDGKEVEQTS GAEBYDAFNKALS LDKKIDLDLQMPKGYDYAASDKSFE
SHERSFVTLYKCFEGTETIDI SFVKNGPLSRADLDTNDTFIVENGSEGLWVWVG
KKATQKERQSAIKYAMEL INK KKYPNNTPVTKVLEGDESVEFKSLFESWQMS
QEKI TSARLFRVSRNGI FKQVANYEPDDLEEDNIMILDVMDKI YVWIGNQFAERI
ADEAHVDKVAQRFI QEDKSGRFRPNQII KLGQSE DGAFKSYFPKWN

Der f 16_iso2_TI

(SEQ ID NO: 41)

MAAHDKNPDVIPIGHTFFFIWRIKQFELVVPKEDYGKFKGDCYIVACCTENP
TGGHSMKESKPI LN GHGYCHIFWIGSESTKDEAGVAAIKSVELDDFLGGYPVQ
HREIEEFESRQFSSYPKNGI IYKGGYESGPTKMIDELKPSLLHVKGKRRPIVYEC
AHSWKVMNNGDVFILLVFNPFVWVTGKHSNRMERTTAIRVANDLKS ELNRFK
LSSVILEDGKEVEQTS GAEBYDAFNKALS LDKKIDLDLQMPKGYDYAASDKSFE
SHERSFVTLYKCFEGTETIDI SFVKNGPLSRADLDTNDTFIVENGSEGLWVWVG
KKATQKERQSAIKYAMEL INK KKYPNNTPVTKVLEGDESVEFKSLFESWQMS
QEKI TSARLFRVSRNGI FKQVANYEPDDLEEDNIMILDVMDKI YVWIGNQFAERI
ADEAHVDKVAQRFLYKRIKVAVNFDQIRL

Der f 7_iso3

(SEQ ID NO: 69)

MMKFLLLIAAVAFVAVSADPIHYDKIT EEINKAIDDAIAAI EKSETIDPMKVPDHT
DKFERHVGI LDFK GELAMRNI EARGLKQMKRQGDANVKGEEGIVKAHLLIGVH
DDIVSMEDLAYKLGDLHPNTHVISDIQDFVVALSLEI SDEGNI TMTSFEVRQFA
NVVNHIGGLSILDP IFGVLSVLTAI PQD TVRKEMTKVLAPAFKRELEKN

Der f 7_iso3_TI

(SEQ ID NO: 42)

MMKFLLLIAAVAFVAVSADPIHYDKIT EEINKAIDDAIAAI EKSETIDPMKVPDLPI
NSNVMLVFWISKVN

Der f 20_iso1

(SEQ ID NO: 70)

MVDQAVIDKLEAGFQKLQSSAECBSLLKKYLTRNVLDACKGRKTMGATLVD
VVQSGFENLD SGVGLYAPDAESYTLFKELFDPVIEDYHKGPKPTDKHPQTD FGD
VNTLCNVDPNNEFV ISTRVRCGRSLQGYFPNCLTEAQYKEMEEKVKGQLNSF
EGELKGTYYPLLGMDKATQQQLIDDHFLFKEGDRFLQAANACRFWVPGCGIFH
NDNKTFLIWVNEEDHLRI ISMQKGGDLKQVFSRLINGVNHIEKKLPFSRDDRLG
FLTFPCNTLGTTRASVHI KL PKLAADRKKLEEVAGKYNLQVRGTAGEHTESVG
GVYDISNKRMRGLTEYQAVKEMQDGILELIIKIEKSM

Der f 20_iso1_TI

(SEQ ID NO: 43)

MVDQAVIDKLEAGFQKLQSSAECBSLLKKYLTRNVLDACKGRKTMGATLVD
VVQSGFENLD SGVGLYAPDAESYTLFKELFDPVIEDYHKGPKPTDKHPQTD FGD
VNTCVMWIIQIMNLSFQHVYVADHCKVIHLIHA

Der f 20_iso1_TI

(SEQ ID NO: 44)

MVDQAVIDKLEAGFQKLQSSAECBSLLKKYLTRNVLDACKGRKTMGATLVD
VVQSGFENLD SGVGLYAPDAESYTLFKELFDPVIEDYHKGPKPTDKHPQTD FGD
VNTLCNVDPNNEFV ISTRVRCGRSLQGYFPNCLTEAQYKEMEEKVKGQLNSF
EGELKGTYYPLLGMDKATQQQLIDDHFLFKEGDRFLQAANACRFWVPGCGIFH
NDNKTFLIWVNEEDHLRI ISMQKGGDLKQVFSRLINGVNHIEKKLPFSRDDRLG
FLTFPCNTLGTTRASVHI KL PKLAADRKNWKKLANITYKYVVLPVNTPKVLA
VFTISVINVVVWLLNIRPSKRCKMFLN

-continued

Der f 20_iso1_TI

(SEQ ID NO: 45)

MVDQAVIDKLEAGFQKLQSSAECHSLLKKYLTRNVLDACKGRKTMGATLVD
VVQSGFENLDSGVGLYAPDAESYTLFKELFDPVI EDYHKGFKPTDKHPQDFDGF
VNTLCNVDPNNEFVISTRVRCGRSLQGYFPNCLTEAQYKEMEEKVKGQLNSF
EGELKGTYYPLLGMDKATQQQLIDHFLFKEGDRFLQAAANACRFWPVGCIGIFH
NDNKTFLIWVNEEDHLRIISMQKGGDLKQVFSRLINGVNHIEKKLPFSRDDRLG
FLTFPCPTNLGTTIRASVHIKLPKLAADRKKLEEVAGKYNLQVRGTAVNTPKVLAV
FTISVINVVWVLLNIRPSKRCKMVFLLN

Der f 20_iso1_TI

(SEQ ID NO: 46)

MVDQAVIDKLEAGFQKLQSSAECHSLLKKYLTRNVLDACKGRKTMGATLVD
VVQSGFENLDSGVGLYAPDAESYTLFKELFDPVI EDYHKGFKPTDKHPQDFDGF
VNTLCNVDPNNEFVISTRVRCGRSLQGYFPNCLTEAQYKEMEEKVKGQLNSF
EGELKGTYYPLLGMDKATQQQLIDHFLFKEGDRFLQAAANACRFWPVGCIGIFH
NDNKTFLIWVNEEDHLRIISMQKGGDLKQVFSRLINGVNHIEKKLPFSRDDRLG
FLTFPCPTNLGTTIRASVHIKLPKLAADRKKLEEVAGKYNLQVRGTAGEHTESV
GVTISVINVVWVLLNIRPSKRCKMVFLLN

Der f 1_iso3

(SEQ ID NO: 71)

MKFALFVVASLIATVYGQSHQYYHTSGLRNLGGSYYRSAGISGVAGLGGLAYG
TGLGYGTRYGYGSLGGLYGLYGLYGLYQAVLAPAQAVGYVAAAPAVAVQAP
AVSYAAAAPAVQTVAVQAPAVSYAAAAPAVAVQHAQAQVSGPIHAAIESRRTV
EVIDGPSTGDAPVASTVVI GPNVQPINLEFQTQASPLAATQNHVPTAPAEPQOSS
YEEQPDLLRQDIVKPVVQDVHETIVPFRRI TQELKPVQESVHQILPRGQERGFYQ
QQQQVRVAQHVAAPA AVAVQPVVQAAPASAVRVAAPAVAYAAPAVSTVS
AAPAAIGVIGVQPAAGYIGYGAGYGTGYGTGYGAKYGTGYGLTSLIGGGSY
GSSYSVQPASVGTGYGTYTSSDAYPIRKK

Der f 1_iso3_TI

(SEQ ID NO: 47)

MKFALFVVASLIATVYGQSHQYYHTSGLRNLGGSYYRSAGISGVAGLGGLAYG
TGLGYGTRYGYGSLGGLYGLYGLYGLYQAVLAPAQAVGYVAAAPAVAVQAP
AVSYAAAAPAVQTVAVQAPAVSYAAAAPAVAVQHAQAQVSGPIHAAIESRRTV
EVIDGPSTGDAPVASTVVI GPNVQPINLEFQTQASPLAATQNHVPTAPAEPQOSS
YEEQPDLLRQDIVKPVVQDVHETIVPFRRI TQELKPVQESVHQILPRGQERGFYQ
QQQQVRVAQHVAAPA AVAVQPVVQAAPASAVRVAAPAVAYAAPAVSTVS
AAPAAIGVIGVQPAAGYIGYGAGYGTGYGTGYGAKYGTGYGLTSLIGVAVH
MDHHIQYNQPA TELVMVTLPIAVMPTQSEKNKLVLPFSF

Der f 1_iso3_TI

(SEQ ID NO: 48)

MKFALFVVASLIATVYGQSHQYYHTSGLRNLGGSYYRSAGISGVAGLGGLAYG
TGLGYGTRYGYGSLGGLYGLYGLYGLYQAVLAPAQAVGYVAAAPAVAVQAP
AVSYAAAAPAVQTVAVQAPAVSYAAAAPAVAVQHAQAQVSGPIHAAIESRRTV
EVIDGPSTGDAPVASTVVI GPNVQPINLEFQTQASPLAATQNHVPTAPAEPQOSS
YEEQPDLLRQDIVKPVVQDVHETIVPFRRI TQELKPVQESVHQILPRGQERGFYQ
QQQQVRVAQHVAAPA AVAVQPVVQAAPASAVRVAAPAVAYAAPAVSTVS
AAPAAIGVIGVQPAAGYIGYGAGYGTGYGTGYGAKYGTGYGLTSLIGVAVH
MDHHIQYNQPA TELVMVTLPIAVMPTQSEKNKLVLPFSF

Der f 15_iso2

(SEQ ID NO: 72)

MKTIYAILSIMACIGLMNASIKRDHNDYSKNPMRIVCYVGTWSVYHKVDPYTI
DIDPFKCTHLMYGFAKIDEYKYTIQVFDPYQDDNHNSWEKRYERFNNLRLKN
PELTTMISLGGWYEGSEKYSDMAANPTYRQQFIQSVLDFLQYKFDGLDLDWE
YPGSRLGNPKIDKQNYLALVRELKDAFEPHGYLLTAAVSPGKDKIDRAYDIKEL
NKLPDWMNVMTYDYHGGWENFYGHNAPLYKRPDETDDELHTYFNVNYTMHY
YLNNGATRDKLVMGVPPFYGRAWSIEDRSKCLKLGDPAKGMSPPGFISGEEGVLS
YIELCQLFPQKEEWHIQYDEYYNAPYGYNDKIWVGDDLASISCKLAFKELGV
SGVIVWLENDFFKGHCGRPKNPLLNKVNHMINGDEKNSFECLILGPSTTPTPTTT
PTTPT
TTEHTSETPKYTTTVDGHLIKCYKEGDIHPHTNIHKYLVCEFVNGGWWVHIMP
CPPGTIWCQEKLTICIGE

Der f 15_iso2_TI

(SEQ ID NO: 49)

MKTIYAILSIMACIGLMNASIKRDHNDYSKNPMRIVCYVGTWSVYHKVDPYTI
DIDPFKCTHLMYGFAKIDEYKYTIQVFDPYQDDNHNSWEKRYERFNNLRLKN
PELTTMISLGGWYEGSEKYSDMAANPTYRQQFIQSVLDFLQYKFDGLDLDWE
LDDLWVTRKSTNKT IWLWLENLKTLLNLMATC

Der f 20_iso2

(SEQ ID NO: 73)

MVDQATLSKLEAGFQKLQNAQDCHSLLKKYLTRDVLQDKTKKTDMGATLL
DVIQSGVENLDSGVGIYAPDAQSYKTFALFDPI IDDYHKGFKPTDKHPQDFDGF
NIEHFVNDVPKNEYVISTRVRCGRSLKGYFPNMLTEAQYKEMETKVKGQLAT
FEGLKGTYYPLLGMDKATQQQLIDHFLFKEGDRFLQAAANACRYWPVGRGIF
HNDKKTFLMWVNEEDHLRIISMQKGGDLKEVFRGLVAVKVKHIEQKIPFSRDDR

-continued

LGYLTFPCPTNLGTTIRASVHIKLPKLAADRKKLEEVAARYNLQVRGTAGEHTES
VGGIYDISNKRMRGLTEYQAVKEMQDGGIIELIKMEKSL

Der f 20_iso2_TI

(SEQ ID NO: 50)

MVDQATLSKLEAGFQKLQNAQDCHSLKKYLTRDVLQDKTKKTDMGATLL
DVIQSGVENLDSGVGIYAPDAQSYKTFALFDPIIDDYHKGFKPTDKHPQTDG
NIEHFVNVDPKNEYVISTRVRCGRSLKGYFPNPLTEAQYKEMETKVKGLAT
FEGELKGTYYPLLGMDKATQQKLIDDHFLFKEGDRFLQANACRYWVGRGIF
HNDKKTFLMWVNEEDHLRIISMQKGGDLKEVFGRLVKAVKHIEQKIPFSRDDR
LGYLTFPCPTNLGTTIRASVHIKLPKLAADRKKLEEVAARYNLQVRGTAGEHTES
VVMILVTVNDEWVSPNTKLLRKCKMASLN

Der f 20_iso2_TI

(SEQ ID NO: 51)

MVDQATLSKLEAGFQKLQNAQDCHSLKKYLTRDVLQDKTKKTDMGATLL
DVIQSGVENLDSGVGIYAPDAQSYKTFALFDPIIDDYHKGFKPTDKHPQTDG
NIEHFVNVDPKNEYVISTRVRCGRSLKGYFPNPLTEAQYKEMETKVKGLAT
FEGELKGTYYPLLGMDKATQQKLIDDHFLFKEGDRFLQANACRYWVGRGIF
HNDKKTFLMWVNEEDHLRIISMQKGGDLKEVFGRLVKAVKHIEQKIPFSRDDR
LGYLTFPCPTNLGTTIRASVHIKLPKLAADRKKLEEVAARYNLQVRGTAGEHTES
VGGIYDISNDEWVSPNTKLLRKCKMASLN

20

The sequences of TI gap peptides included (in underlined form) in the above TI proteins are the following:

-continued

Der f 1_iso1_TI

(SEQ ID NO: 74)

FWPTVTRLWIPLNRNSSIAHLNTDVTAIQYQASNTSNKMVSLKKEAIHT
LHENNNADDQIRNITVSQTAKFIHQM

Der f 1_iso1_TI

(SEQ ID NO: 75)

SNIMMDEQSFNMTMVINQTIMPSTLSVTEVHKASIGSYETVGIQPGVIA
DTDISKPETTS

Der f 1_iso1_TI

(SEQ ID NO: 76)

MVINQTIMPSTLSVTEVHKASIGSYETVGIQPGVIADTDISKPETTS

Der f 1_iso1_TI

(SEQ ID NO: 77)

IGSYETVGIQPGVIADTDISKPETTS

Der f 1_iso2_TI

(SEQ ID NO: 78)

FWPTVTRLWIPLNRNSSIAHLNTDVTAIQYQASNTSNKMVSLKKEAIHT
LHENNNADDQIRNITVSQTAKFIHQM

Der f 1_iso2_TI

(SEQ ID NO: 79)

SNIMMDEQSFNMTMVINQTIMPSTLSVTEVHKASIGSYETVGIQPGVIA
DTDISKPETTS

Der f 1_iso2_TI

(SEQ ID NO: 80)

MVINQTIMPSTLSVTEVHKASIGSYETVGIQPGVIADTDISKPETTS

Der f 1_iso2_TI

(SEQ ID NO: 81)

IGSYETVGIQPGVIADTDISKPETTS

Der f 23_iso1_TI

(SEQ ID NO: 82)

IFVQIGKLYIKVVQVIQDGMKKN

Der f 11_iso1_TI

(SEQ ID NO: 83)

YVINVINWHVKTRNLQITILPKLNHN

Der f 11_iso1_TI

(SEQ ID NO: 84)

TMKYIKNLEFLMNEYRNLQNSNLLKIC

Der f 15_iso1_TI

(SEQ ID NO: 85)

WIGSILDLDWVTRKSTNKTIWLWLENLKTLLNLMATC

Der f 32_iso1_TI

(SEQ ID NO: 86)

LLKKSLLKHHINIGKK

Der f 25_iso1_TI

(SEQ ID NO: 87)

GPLVLVKQPVHNKHKFKIKNFDNGFLKMFPEIRKLPKQFESFMVVQ

Der f 25_iso1_TI

(SEQ ID NO: 88)

MFHHKLPKQFESFMVVQ

Der f 16_iso1_TI

(SEQ ID NO: 89)

YKRIKVAVNFDQIRL

Der f 26_iso1_TI

(SEQ ID NO: 90)

EISVHPALVKKDLVNLNHPHFIVSYPII

Der f 26_iso1_TI

(SEQ ID NO: 91)

KRWKAMAHNRVNHPRK

Der f 23_iso2_TI

(SEQ ID NO: 92)

IFVQIGKLYIKVVQVIQDGMKKN

Der f 13_iso1_TI

(SEQ ID NO: 93)

KVTINLPKHNSVIKK

Der f 7_iso1_TI

(SEQ ID NO: 94)

LPINSNVMVLFVWISKVN

Der f 27_iso1_TI

(SEQ ID NO: 95)

MILTKFVNQSIQLLLMKLLDKLLVSHQLFISRKLNSVHHINCRKF

Der f 7_iso2_TI

(SEQ ID NO: 96)

LPINSNVMVLFVWISKVN

Der f 29_iso1_TI

(SEQ ID NO: 97)

KRWKAMAHNRVNHPRK

Der f 18_iso1_TI

(SEQ ID NO: 98)

KPCIIVPITYRHLTK

Der f 16_iso2_TI

(SEQ ID NO: 99)

YKRIKVAVNFDQIRL

-continued

Der f 7_iso3_TI (SEQ ID NO: 100)
 LPINSNVMLVFWISKVN

Der f 20_iso1_TI (SEQ ID NO: 101)
 CVMWIQIMNLSFQHVYVVADHCKVIHLIHA

Der f 20_iso1_TI (SEQ ID NO: 102)
 WKKLLANITYKYVVLVFNTPKVLAVFTISVINVVWLLNIRPSKRCKMVF
 LN

Der f 20_iso1_TI (SEQ ID NO: 103)
 VNTPKVLAVFTISVINVVWLLNIRPSKRCKMVF LN

Der f 20_iso1_TI (SEQ ID NO: 104)
 TISVINVVWLLNIRPSKRCKMVF LN

Der f 1_iso3_TI (SEQ ID NO: 105)
 EQVMVLLNTEPDMVSLAV

Der f 1_iso3_TI (SEQ ID NO: 106)
 VAHMDHHIQYNQPATELVMTLPPIAVMPTQSEKNKLVLPFSF

Der f 15_iso2_TI (SEQ ID NO: 107)
 WIGSILDLDDWVTRKSTNKTIWLWLENLKTLLNLMATC

Der f 20_iso2_TI (SEQ ID NO: 108)
 VVSMILVTNDEWVSPNTKLLRKCKMASLN

Der f 20_iso2_TI (SEQ ID NO: 109)
 TNDEWVSPNTKLLRKCKMASLN

In conclusion, we were thus able to identify transcription infidelity (TI) events having an impact on the coding sequences of mite allergens, said TI events giving rise to protein variants which are responsible for triggering mite allergy.

Example 5: Analysis of Peanut Allergens: A Correlation Between Transcription Infidelity (TI) Peptides and Peanut Allergy

A—Sensitization to Peanut

In this experiment, we intraperitoneally injected either the peanut extract (400 µg of protein), or the recombinant canonic peanut protein AraH2 (400 µg) in BalbC mice once per week during 3 weeks in order to induce sensitization to peanut (in groups of 10 mice).

The immunization protocol was carried out according to FIG. 12.

The tested adjuvant was LewisX. A control sample without adjuvant was also tested. Then, we measured the rate of IgE against peanut extract in samples collected at day 33 (see FIG. 7A) and day 49 after the beginning of injections (see FIG. 7B).

Our results clearly show that the peanut extract significantly induces sensitization at day 33, and even twice more at day 49, whereas the recombinant canonic peanut protein AraH2 alone cannot induce sensitization since the level of IgE against peanut extract obtained for AraH2 sensitized mice is comparable to that obtained with controls (as shown in FIG. 7).

B—Production of Recombinant AraH2-TI Proteins and Confirmation of Their Allergenicity

We have analyzed AraH2 RNA sequences of *Arachis hypogaea* by Illumina next-generation sequencing and then adapted it to sequential immunization technique in order to produce recombinant Transcription Infidelity (TI) proteins, i.e., AraH2-TI.

In order to define the positions where Transcription Infidelity (TI) events occur, the reads obtained were aligned and we have selected transcription infidelity (TI) positions of deletions which are located at the level of epitopes of AraH2. These TI deletions in epitopes of AraH2 are present in cascade as shown in FIG. 8.

We were able to identify the following TI proteins in peanut (SEQ ID NO: 121 to 131):

Arah2-ref1-g11-id30 (SEQ ID NO: 121)
 MARQQWELQGD RRCSQLERANLRPCEQHLMRKSNVTRIHMDGTRTALVR
 IRTALVTRTRTDVIRTALVHMIGEALDLLSTKRGVAMS

Arah2-ref1-g11-id31 (SEQ ID NO: 122)
 MARQQWELQGD RRCSQLERANLRPCEQHLMQKIQRDEDSYGRDPYSLVR
 IRTALVTRTRTDVIRTALVHMIGEALDLLSTKRGVAMS

Arah2-ref1-g11-id32 (SEQ ID NO: 123)
 MARQQWELQGD RRCSQLERANLRPCEQHLMQKIQRDEDSYGRDPYSPSRI
 RTALVTRTRTDVIRTALVHMIGEALDLLSTKRGVAMS

Arah2-ref1-g11-id33 (SEQ ID NO: 124)
 MARQQWELQGD RRCSQLERANLRPCEQHLMQKIQRDEDSYGRDPYSPSQD
 PYPSPQDPDRDPYSLVHMIGEALDLLSTKRGVAMS

Arah2-ref1-g11-id34 (SEQ ID NO: 125)
 MARQQWELQGD RRCSQLERANLRPCEQHLMQKIQRDEDSYGRDPYSPSQD
 PYPSPQDPDRDPYSPSPYDRRGAGSLSTKRGVAMS

Arah2-ref1-g12-id35 (SEQ ID NO: 126)
 MARQQWELQGD RRCSQLERANLRPCEQHLMQKIQRDEDSYGRDPYSPSQD
 PYPSPQDPDRDPYSPSPYDRRGAGSSQHQRCCNELNEFENNKGACARHC
 NR

Arah2-ref1-g12-id36 (SEQ ID NO: 127)
 MARQQWELQGD RRCSQLERANLRPCEQHLMQKIQRDEDSYGRDPYSPSQD
 PYPSPQDPDRDPYSPSPYDRRGAGSSQHQRCCNELNEFENNQGACARHC
 NR

Arah2-ref2-g13-id37 (SEQ ID NO: 128)
 MARQQWELQGD RRCSQLERANLRPCEQHLMQKIQRDEDSYERDPYSPSQD
 PYPSPYDRRGAGSSQHQRCCNELNEFENNKGACARHCNRSWRTRAIGCR

Arah2-ref2-g13-id38 (SEQ ID NO: 129)
 MARQQWELQGD RRCSQLERANLRPCEQHLMQKIQRDEDSYERDPYSPSQD
 PYPSPYDRRGAGSSQHQRCCNELNEFENNQGACARHCNRSWRTRAIGCR
 GGNRSNSRSGSGTCLNSAALGHHSVATWTSKVA AETDTKHL SOKKKRKEK
 KIAYI

Arah2-ref2-g13-id39
 (SEQ ID NO: 130)
 MARQQWELQGD~~RR~~CQSLERANLRPCEQHL~~M~~QKIQ~~R~~DEDSYERDPYSPSQD
 PYS~~P~~SPYDRRGAGSSQH~~Q~~ERC~~C~~NELNEFENN~~Q~~RCMCEALQQIMENQSDRLQ
 GRQQEQQ~~S~~RGSSGTCLNSAALGHHSVATWT~~S~~KVAAETDTK~~H~~LSQ~~K~~KKRKEK
KIAYI

Arah2-ref2-g13-id40
 (SEQ ID NO: 131)
 MARQQWELQGD~~RR~~CQSLERANLRPCEQHL~~M~~QKIQ~~R~~DEDSYERDPYSPSQD
 PYS~~P~~SPYDRRGAGSSQH~~Q~~ERC~~C~~NELNEFENN~~Q~~RCMCEALQQIMENQSDRLQ
 GRQQEQQ~~S~~RGSSGTCLNSAALGHHSVATWT~~S~~KVAAETDTK~~H~~LSQ~~K~~KKRKEK
KIAYI

The sequences of the TI gap peptides (comprised and underlined in the above TI peanut proteins of SEQ ID NO: 121 to 131) and represented in FIG. 8 as bold, are the following:

Arah2-ref1-g11-id30
 (SEQ ID NO: 110)
 R~~K~~SNVTRIHMDGTRTALV~~R~~IR~~T~~ALV~~R~~TRTDVIR~~T~~ALV~~H~~MIGEALDLLSTK
 RGVAMS

Arah2-ref1-g11-id31
 (SEQ ID NO: 111)
 LV~~R~~IR~~T~~ALV~~R~~TRTDVIR~~T~~ALV~~H~~MIGEALDLLSTKRGVAMS

Arah2-ref1-g11-id32
 (SEQ ID NO: 112)
 RIR~~T~~ALV~~R~~TRTDVIR~~T~~ALV~~H~~MIGEALDLLSTKRGVAMS

Arah2-ref1-g11-id33
 (SEQ ID NO: 113)
 LV~~H~~MIGEALDLLSTKRGVAMS

Arah2-ref1-g11-id34
 (SEQ ID NO: 114)
 LSTKRGVAMS

Arah2-ref1-g12-id35
 (SEQ ID NO: 115)
 KGACARHCNR

Arah2-ref1-g12-id36
 (SEQ ID NO: 116)
 GACARHCNR

Arah2-ref2-g13-id37
 (SEQ ID NO: 117)
 KGACARHCNRSWRTRAIGCRGGNRSN~~S~~RGSSGTCLNSAALGHHSVATWT
 SKVAAETDTK~~H~~LSQ~~K~~KKRKEK~~K~~IAYI

Arah2-ref2-g13-id38
 (SEQ ID NO: 118)
 GACARHCNRSWRTRAIGCRGGNRSN~~S~~RGSSGTCLNSAALGHHSVATWT
 KVAAETDTK~~H~~LSQ~~K~~KKRKEK~~K~~IAYI

Arah2-ref2-g13-id39
 (SEQ ID NO: 119)
 SRGSSGTCLNSAALGHHSVATWT~~S~~KVAAETDTK~~H~~LSQ~~K~~KKRKEK~~K~~IAYI

Arah2-ref2-g13-id40
 (SEQ ID NO: 120)
 HSVATWT~~S~~KVAAETDTK~~H~~LSQ~~K~~KKRKEK~~K~~IAYI

For comparison, the sequences of the canonical proteins are provided in SEQ ID NO: 132 and 133 below:

Arah2-ref1
 (SEQ ID NO: 132)
 MARQQWELQGD~~RR~~CQSLERANLRPCEQHL~~M~~QKIQ~~R~~DEDSYGRDPYSPSQ
 DPYSPSQD~~P~~DRRD~~P~~YSPSPYDRRGAGSSQH~~Q~~ERC~~C~~NELNEFENN~~Q~~RCMCE
 ALQQIMENQSDRLQGRQQEQQ~~F~~KREL~~R~~NLPQQCGLRAP~~Q~~RC~~D~~LEVESGGR
 DRY

Arah2-ref2
 (SEQ ID NO: 133)
 MARQQWELQGD~~RR~~CQSLERANLRPCEQHL~~M~~QKIQ~~R~~DEDSYERDPYSPSQ
 DPYSPSPYDRRGAGSSQH~~Q~~ERC~~C~~NELNEFENN~~Q~~RCMCEALQQIMENQSDR
 LQGRQQEQQ~~F~~KREL~~R~~NLPQQCGLRAP~~Q~~RC~~D~~L~~D~~VESGGRDRY

Then, we produced the recombinant proteins and measured IgE against variants 36, 38 and 40 (said variants 36, 38 and 40 corresponding to TI proteins of SEQ ID NO: 127, 129 and 131 comprising respectively TI gap peptides of SEQ ID NO: 116, 118 and 120).

We used BalbC mice previously described (see FIG. 9). Therefore, FIG. 9 shows that sensitization to peanut induces IgE reactivity specifically against variant-40 and not against variant-38. Variant-40 contains 2 epitopes (epitopes 8 and 9) that are not present in variant-38.

The recombinant variants of AraH2 are injected in BalbC mice in order to induce sensitization to peanut. A negative control is AraH2 recombinant canonic protein which does not induce sensitization.

This example allows to show that the recombinant canonic AraH2 protein (considered in the art as "allergen") is not allergenic at all. On the contrary, the recombinant AraH2-TI proteins are highly allergenic. In conclusion, only AraH2-TI protein variants which result from Transcription Infidelity (TI) deletion events are at the origin of peanut allergy.

C—Prevalence of Peanut Allergy

The data of the percentages of prevalence of allergy in relation to various peanut allergens are known in the literature (Crit Rev Food Sci Nutr. 2013, Peanut allergens: Sáiz JI, Montealegre C, Marina ml, Garcia-Ruiz C), as summarized in Table 6.

TABLE 6

Prevalence of peanut allergy	
Allergen	Prevalence (%)
Ara h 1	90
Ara h 2	90
Ara h 3	50
Ara h 5	13
Ara h 6	38
Ara h 7	43
Ara h 8	70
Ara h 9	45
Ara h 10	21
Ara h 11	21

In this experiment, we have analyzed mRNA of Ara h 1, Ara h 2, Ara h 3, Ara h 5, Ara h 6, Ara h 7, Ara h 8, Ara h 9, Ara h 10 and Ara h 11 by Illumina next-generation sequencing in order to identify transcription infidelity deletion events. We have then selected the deletions present in the repetitions of A bases, and reported the number of the identified deletions in relation to the transcripts' size.

Results of our analysis and comparison with the known prevalence of peanut allergens (from Table 6), are shown in FIG. 10. This figure clearly demonstrates the existence of a

strong correlation between TI deletion events (i.e., number of gap events affecting repetition of A within ORF per 1000 bases) and the prevalence of the peanut allergens. The correlation between these two variables, prevalence and deletions in the repetitions of A, is very highly significant (rho Spearman=0.68; p-value=0.003).

In conclusion, the allergenicity of Ara h 1, Ara h 2, Ara h 3, Ara h 5, Ara h 6, Ara h 7, Ara h 8, Ara h 9, Ara h 10 and Ara h 11 peanut proteins is strictly correlated with transcription infidelity (TI) events and peanut allergy is thus due to production of TI variants of Ara h 1, Ara h 2, Ara h 3, Ara h 5, Ara h 6, Ara h 7, Ara h 8, Ara h 9, Ara h 10 and Ara h 11, resulting from transcription infidelity gap events.

Example 6: Comparison of TI Events in Strongly Allergenic Peanut and Basically Non-Allergenic Green Beans

This study relates to the comparison of the phenomenon of transcription infidelity (TI) deletion events in peanut (which is strongly allergenic) and in another leguminous plant such as the green beans (being basically non-allergenic).

In order to compare a TI phenomenon in these two plants, which affects transcripts of different size and the expression of which also varies, we used the measurement of the (RNA-DNA difference) RDD rate, which is calculated by identifying sequence variations in RNA sequences based on a comparison to a reference sequence, and expressed as the rate of deletions computed on all transcript positions (as illustrated in Table 7 below).

TABLE 7

The example of calculating of RDD rate by identifying variations in RNA sequences based on a comparison to a Reference sequence, and expressed as the rate of deletions computed on all transcript positions.										
Simplistic example:										
Reference	A	T	T	C	G	C	A	G	C	A
	A	T	T	C	G	C	A	G		
	A	T	T	C	G	C	—	G		
	A	T	T	C	G	C	A	G	C	
	A	T	T	C	G	C	A	G	C	A
	A	T	—	C	G	C	A	G	C	A
	A	T	T	C	G	C	A	G	C	A
RNA-seq reads	A	T	T	C	—	C	A	G	C	A
	A	T	T	C	G	C	—	G	C	A
	A	T	T	C	G	C	A	G	C	A
	A	T	—	C	G	C	—	G	C	A
	A	T	T	C	G	C	A	G	C	A
	A	T	T	C	G	C	A	G	C	A
			T	C	G	C	A	G	C	A
				C	G	C	A	G	C	A
	↓	↓	↓	↓	↓	↓	↓	↓	↓	↓
Number of RDD	0	0	2	0	1	0	3	0	0	0
Number of reads	12	12	13	14	14	14	14	14	12	11
RDD rate =	$\text{RDD rate} = \frac{0+0+2+0+1+0+3+0+0+0}{12+12+13+14+14+14+14+14+14+12+11} = \frac{6}{130}$									

Our results of comparing RDD rate in peanuts and in green beans clearly show that the total RDD rate corresponding to TI deletion events, is significantly higher in peanuts than in green beans, and especially in relation to TI deletion events present in coding regions (ORF) affecting the repetitions of A or T bases. These differences are very

highly significant (test of wilcoxon p-value<0.001) as demonstrated in FIG. 11.

The above data further confirm that TI deletion events which are highly present in allergenic peanut proteins, are necessarily at the origin of peanut allergy.

 SEQUENCE LISTING

<160> NUMBER OF SEQ ID NOS: 133

<210> SEQ ID NO 1
 <211> LENGTH: 16
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: TI peptide

<400> SEQUENCE: 1

Leu Trp His Leu Phe Gln Lys Cys Leu Glu Arg Arg Arg Ser Met Asn
 1 5 10 15

<210> SEQ ID NO 2
 <211> LENGTH: 53
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: TI peptide

<400> SEQUENCE: 2

Leu Pro Ala Phe Trp Leu Leu Pro Leu Gln Arg Ile Arg Trp Asn Met
 1 5 10 15

Ser Pro Pro Val Arg Asn Leu Ser Ser Pro Arg Lys His Ile Ser Arg
 20 25 30

Lys Arg Ile Trp Pro Leu Ile Pro Ala Arg Arg Thr Phe Ala Pro His
 35 40 45

Ser Ala Arg Lys Leu
 50

<210> SEQ ID NO 3
 <211> LENGTH: 44
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: TI peptide

<400> SEQUENCE: 3

Glu Ala Phe Gln Ala Val Arg Asn Leu Leu His Ala Ser Ile Arg Lys
 1 5 10 15

Leu Arg Ser Phe Arg Val Arg Asn Ser Ser Lys Gln Arg Met Asn Ser
 20 25 30

Arg Ile Lys Ser Thr Pro Leu Pro Arg His Ser Leu
 35 40

<210> SEQ ID NO 4
 <211> LENGTH: 25
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: TI peptide

<400> SEQUENCE: 4

Phe Trp Val Pro Arg Ser Lys Thr Lys Asn Asn Gln Tyr Ala Val Arg
 1 5 10 15

Lys Met Lys Asp Ser Ser Val Thr Lys
 20 25

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<210> SEQ ID NO 5
 <211> LENGTH: 64
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: TI peptide

<400> SEQUENCE: 5

Lys Ser Thr Cys Ser Ser Ala Trp Arg Thr Val Leu Ser Pro Ser Lys
 1 5 10 15
 Ala Trp Pro Ala Ser Ala Trp Ser Gly Pro Arg Arg Trp Thr Thr Arg
 20 25 30
 Pro Trp Arg Asn Ser Thr Lys Pro Ser Arg Pro Cys Pro Cys Thr Ser
 35 40 45
 Gly Cys Pro Ser Thr Gln Pro Ser Trp Arg Ser Ser Ala Thr Ser Arg
 50 55 60

<210> SEQ ID NO 6
 <211> LENGTH: 54
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: TI protein

<400> SEQUENCE: 6

Met Lys Leu Leu Ile Leu Thr Cys Leu Val Ala Val Ala Leu Ala Arg
 1 5 10 15
 Pro Lys His Pro Ile Lys His Gln Gly Leu Pro Gln Glu Val Leu Asn
 20 25 30
 Glu Asn Leu Leu Arg Phe Leu Trp His Leu Phe Gln Lys Cys Leu Glu
 35 40 45
 Arg Arg Arg Ser Met Asn
 50

<210> SEQ ID NO 7
 <211> LENGTH: 58
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: TI protein

<400> SEQUENCE: 7

Met Lys Phe Phe Ile Leu Pro Ala Phe Trp Leu Leu Pro Leu Gln Arg
 1 5 10 15
 Ile Arg Trp Asn Met Ser Pro Pro Val Arg Asn Leu Ser Ser Pro Arg
 20 25 30
 Lys His Ile Ser Arg Lys Arg Ile Trp Pro Leu Ile Pro Ala Arg Arg
 35 40 45
 Thr Phe Ala Pro His Ser Ala Arg Lys Leu
 50 55

<210> SEQ ID NO 8
 <211> LENGTH: 107
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: TI protein

<400> SEQUENCE: 8

Met Pro Leu Asn Thr Ile Tyr Lys Gln Pro Gln Asn Gln Ile Ile Ile
 1 5 10 15

-continued

His Ser Ala Pro Pro Ser Leu Leu Val Leu Tyr Phe Gly Lys Lys Glu
 20 25 30

Leu Arg Ala Met Lys Val Leu Ile Leu Ala Cys Leu Val Ala Leu Ala
 35 40 45

Leu Ala Arg Glu Leu Glu Glu Leu Asn Val Pro Gly Glu Ile Val Glu
 50 55 60

Ala Phe Gln Ala Val Arg Asn Leu Leu His Ala Ser Ile Arg Lys Leu
 65 70 75 80

Arg Ser Phe Arg Val Arg Asn Ser Ser Lys Gln Arg Met Asn Ser Arg
 85 90 95

Ile Lys Ser Thr Pro Leu Pro Arg His Ser Leu
 100 105

<210> SEQ ID NO 9
 <211> LENGTH: 42
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: TI protein

<400> SEQUENCE: 9

Met Met Lys Ser Phe Phe Leu Val Val Thr Ile Leu Ala Leu Thr Leu
 1 5 10 15

Pro Phe Trp Val Pro Arg Ser Lys Thr Lys Asn Asn Gln Tyr Ala Val
 20 25 30

Arg Lys Met Lys Asp Ser Ser Val Thr Lys
 35 40

<210> SEQ ID NO 10
 <211> LENGTH: 179
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: TI protein

<400> SEQUENCE: 10

Met Lys Cys Leu Leu Leu Ala Leu Ala Leu Thr Cys Gly Ala Gln Ala
 1 5 10 15

Leu Ile Val Thr Gln Thr Met Lys Gly Leu Asp Ile Gln Lys Val Ala
 20 25 30

Gly Thr Trp Tyr Ser Leu Ala Met Ala Ala Ser Asp Ile Ser Leu Leu
 35 40 45

Asp Ala Gln Ser Ala Pro Leu Arg Val Tyr Val Glu Glu Leu Lys Pro
 50 55 60

Thr Pro Glu Gly Asp Leu Glu Ile Leu Leu Gln Lys Trp Glu Asn Gly
 65 70 75 80

Glu Cys Ala Gln Lys Lys Ile Ile Ala Glu Lys Thr Lys Ile Pro Ala
 85 90 95

Val Phe Lys Ile Asp Ala Leu Asn Glu Asn Lys Val Leu Val Leu Asp
 100 105 110

Thr Asp Tyr Lys Ser Thr Cys Ser Ser Ala Trp Arg Thr Val Leu Ser
 115 120 125

Pro Ser Lys Ala Trp Pro Ala Ser Ala Trp Ser Gly Pro Arg Arg Trp
 130 135 140

Thr Thr Arg Pro Trp Arg Asn Ser Thr Lys Pro Ser Arg Pro Cys Pro
 145 150 155 160

Cys Thr Ser Gly Cys Pro Ser Thr Gln Pro Ser Trp Arg Ser Ser Ala

-continued

165 170 175

Thr Ser Arg

<210> SEQ ID NO 11
 <211> LENGTH: 200
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: CSN1S1 protein

<400> SEQUENCE: 11

Met Lys Leu Leu Ile Leu Thr Cys Leu Val Ala Val Ala Leu Ala Arg
 1 5 10 15

Pro Lys His Pro Ile Lys His Gln Gly Leu Pro Gln Glu Val Leu Asn
 20 25 30

Glu Asn Leu Leu Arg Phe Phe Val Ala Pro Phe Pro Glu Val Phe Gly
 35 40 45

Lys Glu Lys Val Asn Glu Leu Ser Lys Asp Ile Gly Ser Glu Ser Thr
 50 55 60

Glu Asp Gln Ala Met Glu Asp Ile Lys Gln Met Glu Ala Glu Ser Ile
 65 70 75 80

Ser Ser Ser Glu Glu Ile Val Pro Asn Ser Val Glu Gln Lys His Ile
 85 90 95

Gln Lys Glu Asp Val Pro Ser Glu Arg Tyr Leu Gly Tyr Leu Glu Ile
 100 105 110

Val Pro Asn Ser Ala Glu Glu Arg Leu His Ser Met Lys Glu Gly Ile
 115 120 125

His Ala Gln Gln Lys Glu Pro Met Ile Gly Val Asn Gln Glu Leu Ala
 130 135 140

Tyr Phe Tyr Pro Glu Leu Phe Arg Gln Phe Tyr Gln Leu Asp Ala Tyr
 145 150 155 160

Pro Ser Gly Ala Trp Tyr Tyr Val Pro Leu Gly Thr Gln Tyr Thr Asp
 165 170 175

Ala Pro Ser Phe Ser Asp Ile Pro Asn Pro Ile Gly Ser Glu Asn Ser
 180 185 190

Glu Lys Thr Thr Met Pro Leu Trp
 195 200

<210> SEQ ID NO 12
 <211> LENGTH: 222
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: CSN1S2 protein

<400> SEQUENCE: 12

Met Lys Phe Phe Ile Phe Thr Cys Leu Leu Ala Val Ala Leu Ala Lys
 1 5 10 15

Asn Thr Met Glu His Val Ser Ser Ser Glu Glu Ser Ile Ile Ser Gln
 20 25 30

Glu Thr Tyr Lys Gln Glu Lys Asn Met Ala Ile Asn Pro Ser Lys Glu
 35 40 45

Asn Leu Cys Ser Thr Phe Cys Lys Glu Val Val Arg Asn Ala Asn Glu
 50 55 60

Glu Glu Tyr Ser Ile Gly Ser Ser Ser Glu Glu Ser Ala Glu Val Ala
 65 70 75 80

Thr Glu Glu Val Lys Ile Thr Val Asp Asp Lys His Tyr Gln Lys Ala

-continued

<400> SEQUENCE: 17

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Met Lys Phe Val Leu Ala Ile Ala Ser Leu Leu Val Leu Ser Thr Val
1           5           10           15
Tyr Ala Arg Pro Ala Ser Ile Lys Thr Phe Glu Glu Phe Lys Lys Ala
20           25           30
Phe Asn Lys Asn Tyr Ala Thr Val Glu Glu Glu Glu Val Ala Arg Lys
35           40           45
Asn Phe Leu Glu Ser Leu Lys Tyr Val Glu Ala Asn Lys Gly Ala Ile
50           55           60
Asn His Leu Ser Asp Leu Ser Leu Asp Glu Phe Lys Asn Arg Tyr Leu
65           70           75           80
Met Ser Ala Glu Ala Phe Glu Gln Leu Lys Thr Gln Phe Asp Leu Asn
85           90           95
Ala Glu Thr Ser Ala Cys Arg Ile Asn Ser Val Asn Val Pro Ser Glu
100          105          110
Leu Asp Leu Arg Ser Leu Arg Thr Val Thr Pro Ile Arg Met Gln Gly
115          120          125
Gly Cys Gly Ser Cys Trp Ala Phe Ser Gly Val Ala Ala Thr Glu Ser
130          135          140
Ala Tyr Leu Ala Tyr Arg Asn Thr Ser Leu Asp Leu Ser Glu Gln Glu
145          150          155          160
Leu Val Asp Cys Ala Ser Gln His Gly Cys His Gly Asp Thr Ile Pro
165          170          175
Arg Gly Ile Glu Tyr Ile Gln Gln Asn Gly Val Val Glu Glu Arg Ser
180          185          190
Tyr Pro Tyr Val Ala Arg Glu Gln Gln Cys Arg Arg Pro Asn Ser Gln
195          200          205
His Tyr Gly Ile Ser Asn Tyr Cys Gln Ile Tyr Pro Pro Asp Val Lys
210          215          220
Gln Ile Arg Glu Ala Leu Thr Gln Thr His Thr Ala Ile Ala Val Ile
225          230          235          240
Ile Gly Ile Lys Asp Leu Arg Ala Ser Asn Ile Met Met Asp Glu Gln
245          250          255
Ser Phe Asn Met Thr Met Val Ile Asn Gln Thr Ile Met Pro Ser Thr
260          265          270
Leu Ser Val Thr Glu Val His Lys Ala Ser Ile Ile Gly Ser Tyr Glu
275          280          285
Thr Val Gly Ile Gln Pro Gly Val Ile Ala Asp Thr Asp Ile Ser Lys
290          295          300
Pro Glu Thr Thr Ser
305

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<210> SEQ ID NO 18

<211> LENGTH: 309

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: TI protein (Der f 1_iso1_TI)

<400> SEQUENCE: 18

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Met Lys Phe Val Leu Ala Ile Ala Ser Leu Leu Val Leu Ser Thr Val
1           5           10           15
Tyr Ala Arg Pro Ala Ser Ile Lys Thr Phe Glu Glu Phe Lys Lys Ala
20           25           30
Phe Asn Lys Asn Tyr Ala Thr Val Glu Glu Glu Glu Val Ala Arg Lys

```


-continued

85					90					95					
Ala	Glu	Thr	Ser	Ala	Cys	Arg	Ile	Asn	Ser	Val	Asn	Val	Pro	Ser	Glu
	100							105					110		
Leu	Asp	Leu	Arg	Ser	Leu	Arg	Thr	Val	Thr	Pro	Ile	Arg	Met	Gln	Gly
	115						120					125			
Gly	Cys	Gly	Ser	Cys	Trp	Ala	Phe	Ser	Gly	Val	Ala	Ala	Thr	Glu	Ser
	130					135					140				
Ala	Tyr	Leu	Ala	Tyr	Arg	Asn	Thr	Ser	Leu	Asp	Leu	Ser	Glu	Gln	Glu
	145				150					155				160	
Leu	Val	Asp	Cys	Ala	Ser	Gln	His	Gly	Cys	His	Gly	Asp	Thr	Ile	Pro
			165						170					175	
Arg	Gly	Ile	Glu	Tyr	Ile	Gln	Gln	Asn	Gly	Val	Val	Glu	Glu	Arg	Ser
		180						185					190		
Tyr	Pro	Tyr	Val	Ala	Arg	Glu	Gln	Gln	Cys	Arg	Arg	Pro	Asn	Ser	Gln
		195				200						205			
His	Tyr	Gly	Ile	Ser	Asn	Tyr	Cys	Gln	Ile	Tyr	Pro	Pro	Asp	Val	Lys
	210				215						220				
Gln	Ile	Arg	Glu	Ala	Leu	Thr	Gln	Thr	His	Thr	Ala	Ile	Ala	Val	Ile
	225				230					235					240
Ile	Gly	Ile	Lys	Asp	Leu	Arg	Ala	Phe	Gln	His	Tyr	Asp	Gly	Arg	Thr
			245						250					255	
Ile	Ile	Gln	His	Asp	Asn	Gly	Tyr	Gln	Pro	Asn	Tyr	His	Ala	Val	Asn
		260						265					270		
Ile	Val	Gly	Tyr	Gly	Ser	Thr	Gln	Gly	Val	Asp	Ile	Gly	Ser	Tyr	Glu
		275					280					285			
Thr	Val	Gly	Ile	Gln	Pro	Gly	Val	Ile	Ala	Asp	Thr	Asp	Ile	Ser	Lys
	290					295					300				
Pro	Glu	Thr	Thr	Ser											
	305														

<210> SEQ ID NO 20

<211> LENGTH: 222

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: TI protein (Der f 1_iso2_TI)

<400> SEQUENCE: 20

Met	Lys	Phe	Val	Leu	Ala	Ile	Ala	Ser	Leu	Leu	Val	Leu	Ser	Thr	Val
1				5					10					15	
Tyr	Ala	Arg	Pro	Ala	Ser	Ile	Lys	Thr	Phe	Glu	Glu	Phe	Lys	Lys	Ala
			20					25					30		
Phe	Asn	Lys	Asn	Tyr	Ala	Thr	Val	Glu	Glu	Glu	Glu	Val	Ala	Arg	Lys
		35					40					45			
Asn	Phe	Leu	Glu	Ser	Leu	Lys	Tyr	Val	Glu	Ala	Asn	Lys	Gly	Ala	Ile
	50					55					60				
Asn	His	Leu	Ser	Asp	Leu	Ser	Leu	Asp	Glu	Phe	Lys	Asn	Arg	Tyr	Leu
	65				70					75				80	
Met	Ser	Ala	Glu	Ala	Phe	Glu	Gln	Leu	Lys	Thr	Gln	Phe	Asp	Leu	Asn
			85						90					95	
Ala	Glu	Thr	Ser	Ala	Cys	Arg	Ile	Asn	Ser	Val	Asn	Val	Pro	Ser	Glu
			100					105					110		
Leu	Asp	Leu	Arg	Ser	Leu	Arg	Thr	Val	Thr	Pro	Ile	Arg	Met	Gln	Gly
		115					120					125			
Gly	Cys	Gly	Ser	Cys	Trp	Ala	Phe	Ser	Gly	Val	Ala	Ala	Thr	Glu	Ser

-continued

130 135 140

Ala Phe Trp Pro Thr Val Thr Arg Leu Trp Ile Phe Leu Asn Arg Asn
 145 150 155 160

Ser Ser Ile Ala His Leu Asn Thr Asp Val Thr Ala Ile Gln Tyr Gln
 165 170 175

Glu Ala Ser Asn Thr Ser Asn Lys Met Val Ser Leu Lys Lys Glu Ala
 180 185 190

Ile His Thr Leu His Glu Asn Asn Asn Ala Asp Asp Gln Ile Arg Asn
 195 200 205

Ile Thr Val Ser Gln Thr Thr Ala Lys Phe Ile His Gln Met
 210 215 220

<210> SEQ ID NO 21
 <211> LENGTH: 309
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: TI protein (Der f 1_iso2_TI)

<400> SEQUENCE: 21

Met Lys Phe Val Leu Ala Ile Ala Ser Leu Leu Val Leu Ser Thr Val
 1 5 10 15

Tyr Ala Arg Pro Ala Ser Ile Lys Thr Phe Glu Glu Phe Lys Lys Ala
 20 25 30

Phe Asn Lys Asn Tyr Ala Thr Val Glu Glu Glu Glu Val Ala Arg Lys
 35 40 45

Asn Phe Leu Glu Ser Leu Lys Tyr Val Glu Ala Asn Lys Gly Ala Ile
 50 55 60

Asn His Leu Ser Asp Leu Ser Leu Asp Glu Phe Lys Asn Arg Tyr Leu
 65 70 75 80

Met Ser Ala Glu Ala Phe Glu Gln Leu Lys Thr Gln Phe Asp Leu Asn
 85 90 95

Ala Glu Thr Ser Ala Cys Arg Ile Asn Ser Val Asn Val Pro Ser Glu
 100 105 110

Leu Asp Leu Arg Ser Leu Arg Thr Val Thr Pro Ile Arg Met Gln Gly
 115 120 125

Gly Cys Gly Ser Cys Trp Ala Phe Ser Gly Val Ala Ala Thr Glu Ser
 130 135 140

Ala Tyr Leu Ala Tyr Arg Asn Thr Ser Leu Asp Leu Ser Glu Gln Glu
 145 150 155 160

Leu Val Asp Cys Ala Ser Gln His Gly Cys His Gly Asp Thr Ile Pro
 165 170 175

Arg Gly Ile Glu Tyr Ile Gln Gln Asn Gly Val Val Glu Glu Arg Ser
 180 185 190

Tyr Pro Tyr Val Ala Arg Glu Gln Gln Cys Arg Arg Pro Asn Ser Gln
 195 200 205

His Tyr Gly Ile Ser Asn Tyr Cys Gln Ile Tyr Pro Pro Asp Val Lys
 210 215 220

Gln Ile Arg Glu Ala Leu Thr Gln Thr His Thr Ala Ile Ala Val Ile
 225 230 235 240

Ile Gly Ile Lys Asp Leu Arg Ala Ser Asn Ile Met Met Asp Glu Gln
 245 250 255

Ser Phe Asn Met Thr Met Val Ile Asn Gln Thr Ile Met Pro Ser Thr
 260 265 270

Leu Ser Val Thr Glu Val His Lys Ala Ser Ile Ile Gly Ser Tyr Glu

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      275              280              285
Thr Val Gly Ile Gln Pro Gly Val Ile Ala Asp Thr Asp Ile Ser Lys
  290              295              300

Pro Glu Thr Thr Ser
305

<210> SEQ ID NO 22
<211> LENGTH: 309
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: TI protein (Der f 1_iso2_TI )

<400> SEQUENCE: 22
Met Lys Phe Val Leu Ala Ile Ala Ser Leu Leu Val Leu Ser Thr Val
 1              5              10              15
Tyr Ala Arg Pro Ala Ser Ile Lys Thr Phe Glu Glu Phe Lys Lys Ala
 20              25              30
Phe Asn Lys Asn Tyr Ala Thr Val Glu Glu Glu Glu Val Ala Arg Lys
 35              40              45
Asn Phe Leu Glu Ser Leu Lys Tyr Val Glu Ala Asn Lys Gly Ala Ile
 50              55              60
Asn His Leu Ser Asp Leu Ser Leu Asp Glu Phe Lys Asn Arg Tyr Leu
 65              70              75              80
Met Ser Ala Glu Ala Phe Glu Gln Leu Lys Thr Gln Phe Asp Leu Asn
 85              90              95
Ala Glu Thr Ser Ala Cys Arg Ile Asn Ser Val Asn Val Pro Ser Glu
 100             105             110
Leu Asp Leu Arg Ser Leu Arg Thr Val Thr Pro Ile Arg Met Gln Gly
 115             120             125
Gly Cys Gly Ser Cys Trp Ala Phe Ser Gly Val Ala Ala Thr Glu Ser
 130             135             140
Ala Tyr Leu Ala Tyr Arg Asn Thr Ser Leu Asp Leu Ser Glu Gln Glu
 145             150             155             160
Leu Val Asp Cys Ala Ser Gln His Gly Cys His Gly Asp Thr Ile Pro
 165             170             175
Arg Gly Ile Glu Tyr Ile Gln Gln Asn Gly Val Val Glu Glu Arg Ser
 180             185             190
Tyr Pro Tyr Val Ala Arg Glu Gln Gln Cys Arg Arg Pro Asn Ser Gln
 195             200             205
His Tyr Gly Ile Ser Asn Tyr Cys Gln Ile Tyr Pro Pro Asp Val Lys
 210             215             220
Gln Ile Arg Glu Ala Leu Thr Gln Thr His Thr Ala Ile Ala Val Ile
 225             230             235             240
Ile Gly Ile Lys Asp Leu Arg Ala Phe Gln His Tyr Asp Gly Arg Thr
 245             250             255
Ile Ile Gln His Glu Met Val Ile Asn Gln Thr Ile Met Pro Ser Thr
 260             265             270
Leu Ser Val Thr Glu Val His Lys Ala Ser Ile Ile Gly Ser Tyr Glu
 275             280             285
Thr Val Gly Ile Gln Pro Gly Val Ile Ala Asp Thr Asp Ile Ser Lys
 290             295             300

Pro Glu Thr Thr Ser
305

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<210> SEQ ID NO 23
<211> LENGTH: 309
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 1_iso2_TI

<400> SEQUENCE: 23

Met Lys Phe Val Leu Ala Ile Ala Ser Leu Leu Val Leu Ser Thr Val
1          5          10          15

Tyr Ala Arg Pro Ala Ser Ile Lys Thr Phe Glu Glu Phe Lys Lys Ala
20          25          30

Phe Asn Lys Asn Tyr Ala Thr Val Glu Glu Glu Glu Val Ala Arg Lys
35          40          45

Asn Phe Leu Glu Ser Leu Lys Tyr Val Glu Ala Asn Lys Gly Ala Ile
50          55          60

Asn His Leu Ser Asp Leu Ser Leu Asp Glu Phe Lys Asn Arg Tyr Leu
65          70          75          80

Met Ser Ala Glu Ala Phe Glu Gln Leu Lys Thr Gln Phe Asp Leu Asn
85          90          95

Ala Glu Thr Ser Ala Cys Arg Ile Asn Ser Val Asn Val Pro Ser Glu
100         105         110

Leu Asp Leu Arg Ser Leu Arg Thr Val Thr Pro Ile Arg Met Gln Gly
115         120         125

Gly Cys Gly Ser Cys Trp Ala Phe Ser Gly Val Ala Ala Thr Glu Ser
130         135         140

Ala Tyr Leu Ala Tyr Arg Asn Thr Ser Leu Asp Leu Ser Glu Gln Glu
145         150         155         160

Leu Val Asp Cys Ala Ser Gln His Gly Cys His Gly Asp Thr Ile Pro
165         170         175

Arg Gly Ile Glu Tyr Ile Gln Gln Asn Gly Val Val Glu Glu Arg Ser
180         185         190

Tyr Pro Tyr Val Ala Arg Glu Gln Gln Cys Arg Arg Pro Asn Ser Gln
195         200         205

His Tyr Gly Ile Ser Asn Tyr Cys Gln Ile Tyr Pro Pro Asp Val Lys
210         215         220

Gln Ile Arg Glu Ala Leu Thr Gln Thr His Thr Ala Ile Ala Val Ile
225         230         235         240

Ile Gly Ile Lys Asp Leu Arg Ala Phe Gln His Tyr Asp Gly Arg Thr
245         250         255

Ile Ile Gln His Asp Asn Gly Tyr Gln Pro Asn Tyr His Ala Val Asn
260         265         270

Ile Val Gly Tyr Gly Ser Thr Gln Gly Val Asp Ile Gly Ser Tyr Glu
275         280         285

Thr Val Gly Ile Gln Pro Gly Val Ile Ala Asp Thr Asp Ile Ser Lys
290         295         300

Pro Glu Thr Thr Ser
305

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<210> SEQ ID NO 24
<211> LENGTH: 177
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 23_iso1_TI

<400> SEQUENCE: 24

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Met Lys Phe Asn Ile Thr Ile Ala Phe Val Ser Leu Ala Ile Leu Ile
 1 5 10 15
 His Ser Ser Tyr Ala Asp Ile Asp His Phe Asp Asn Asp Asp Gln Asn
 20 25 30
 Ser Ser Thr Ser Arg Pro Asp Asp Asp Pro Thr Thr Met Ile Asp Val
 35 40 45
 Gln Thr Thr Thr Val Gln Pro Ser Ser Met Pro Thr Thr Ser Glu Ser
 50 55 60
 Gln Ser Thr Val Lys Pro Thr Thr Thr Thr Val Lys Pro Ser Pro Thr
 65 70 75 80
 Thr Val Lys Leu Thr Thr Thr Thr Val Lys Pro Thr Thr Thr Thr Val
 85 90 95
 Lys Pro Thr Thr Thr Thr Val Lys Pro Ser Pro Thr Thr Val Lys Pro
 100 105 110
 Thr Thr Thr Thr Val Lys Pro Ser Pro Thr Thr Thr Thr Thr Thr
 115 120 125
 Thr Glu Gln Pro Glu Asp Glu Phe Glu Cys Pro Thr Arg Phe Gly Tyr
 130 135 140
 Phe Ala Asp Pro Lys Asp Pro Cys Lys Phe Ile Phe Val Gln Ile Gly
 145 150 155 160
 Lys Leu Tyr Ile Lys Val Val Gln Val Ile Gln Asp Gly Met Lys Lys
 165 170 175

Asn

<210> SEQ ID NO 25
 <211> LENGTH: 439
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 11_isol_TI

<400> SEQUENCE: 25

Met Ser Ala Arg Thr Ala Lys Tyr Met Tyr Arg Ser Ser Gly Ala Gly
 1 5 10 15
 Ala Ser Gly Asp Ile Ser Val Glu Tyr Gly Thr Asp Leu Gly Ala Leu
 20 25 30
 Thr Arg Leu Glu Asp Lys Ile Arg Leu Leu Ser Asp Asp Leu Glu Ser
 35 40 45
 Glu Arg Glu Met Arg Gln Arg Ile Glu Arg Glu Lys Ala Glu Leu Gln
 50 55 60
 Ile Gln Val Met Ser Leu Gly Glu Arg Leu Glu Glu Ala Glu Gly Ser
 65 70 75 80
 Ser Glu Ser Val Thr Glu Met Asn Lys Lys Arg Asp Ser Glu Leu Ala
 85 90 95
 Lys Leu Arg Lys Leu Leu Glu Asp Val His Ile Glu Ser Glu Glu Thr
 100 105 110
 Ala His His Leu Arg Gln Lys His Gln Ala Ala Ile Gln Glu Met Gln
 115 120 125
 Asp Gln Leu Asp Gln Leu Gln Lys Ala Lys Asn Lys Ser Asp Lys Glu
 130 135 140
 Lys Gln Lys Phe Gln Ala Glu Val Phe Glu Leu Leu Ala Gln Leu Glu
 145 150 155 160
 Thr Ala Asn Lys Glu Lys Leu Thr Ala Leu Lys Asn Val Glu Lys Leu
 165 170 175
 Glu Tyr Thr Val His Glu Leu Asn Ile Lys Ile Glu Glu Ile Asn Arg

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180	185	190
Thr Val Ile Glu Leu Thr Ser His Lys Gln Arg Leu Ser Gln Glu Asn		
195	200	205
Thr Glu Leu Ile Lys Glu Val His Glu Val Lys Leu Gln Leu Asp Asn		
210	215	220
Ala Asn His Leu Lys Thr Gln Ile Ala Gln Gln Leu Glu Asp Thr Arg		
225	230	235
240		
His Arg Leu Glu Glu Glu Glu Arg Lys Arg Ala Ser Leu Glu Asn His		
245	250	255
Ala His Thr Leu Glu Val Glu Leu Glu Ser Leu Lys Val Gln Leu Asp		
260	265	270
Glu Glu Ser Glu Ala Arg Leu Glu Leu Glu Arg Gln Leu Thr Lys Ala		
275	280	285
Asn Gly Asp Ala Ala Ser Trp Lys Ser Lys Tyr Glu Ala Glu Leu Gln		
290	295	300
Ala His Ala Asp Glu Val Glu Glu Leu Arg Arg Lys Met Ala Gln Lys		
305	310	315
320		
Ile Ser Glu Tyr Glu Glu Gln Leu Glu Ala Leu Leu Asn Lys Cys Ser		
325	330	335
Ser Leu Glu Lys Gln Lys Ser Arg Leu Gln Ser Glu Val Glu Val Leu		
340	345	350
Ile Met Asp Leu Glu Lys Ala Thr Ala His Ala Gln Gln Leu Glu Lys		
355	360	365
Arg Val Ala Gln Leu Glu Lys Ile Asn Leu Asp Leu Lys Asn Lys Leu		
370	375	380
Glu Glu Val Thr Met Leu Met Glu Gln Ala Gln Lys Glu Leu Arg Val		
385	390	395
400		
Lys Ile Ala Glu Leu Gln Lys Leu Gln His Glu Tyr Glu Asn Tyr Val		
405	410	415
Ile Asn Val Ile Asn Trp His Val Lys Thr Arg Asn Leu Gln Thr Ile		
420	425	430
Leu Pro Lys Leu Asn His Asn		
435		

<210> SEQ ID NO 26
 <211> LENGTH: 694
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 11_isol_TI

<400> SEQUENCE: 26

Met Ser Ala Arg Thr Ala Lys Tyr Met Tyr Arg Ser Ser Gly Ala Gly
1 5 10 15
Ala Ser Gly Asp Ile Ser Val Glu Tyr Gly Thr Asp Leu Gly Ala Leu
20 25 30
Thr Arg Leu Glu Asp Lys Ile Arg Leu Leu Ser Asp Asp Leu Glu Ser
35 40 45
Glu Arg Glu Met Arg Gln Arg Ile Glu Arg Glu Lys Ala Glu Leu Gln
50 55 60
Ile Gln Val Met Ser Leu Gly Glu Arg Leu Glu Glu Ala Glu Gly Ser
65 70 75 80
Ser Glu Ser Val Thr Glu Met Asn Lys Lys Arg Asp Ser Glu Leu Ala
85 90 95
Lys Leu Arg Lys Leu Leu Glu Asp Val His Ile Glu Ser Glu Glu Thr

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100					105					110					
Ala	His	His	Leu	Arg	Gln	Lys	His	Gln	Ala	Ala	Ile	Gln	Glu	Met	Gln
	115					120					125				
Asp	Gln	Leu	Asp	Gln	Leu	Gln	Lys	Ala	Lys	Asn	Lys	Ser	Asp	Lys	Glu
	130					135					140				
Lys	Gln	Lys	Phe	Gln	Ala	Glu	Val	Phe	Glu	Leu	Leu	Ala	Gln	Leu	Glu
	145					150					155				160
Thr	Ala	Asn	Lys	Glu	Lys	Leu	Thr	Ala	Leu	Lys	Asn	Val	Glu	Lys	Leu
				165					170					175	
Glu	Tyr	Thr	Val	His	Glu	Leu	Asn	Ile	Lys	Ile	Glu	Glu	Ile	Asn	Arg
			180					185						190	
Thr	Val	Ile	Glu	Leu	Thr	Ser	His	Lys	Gln	Arg	Leu	Ser	Gln	Glu	Asn
			195					200					205		
Thr	Glu	Leu	Ile	Lys	Glu	Val	His	Glu	Val	Lys	Leu	Gln	Leu	Asp	Asn
	210					215					220				
Ala	Asn	His	Leu	Lys	Thr	Gln	Ile	Ala	Gln	Gln	Leu	Glu	Asp	Thr	Arg
	225					230					235				240
His	Arg	Leu	Glu	Glu	Glu	Glu	Arg	Lys	Arg	Ala	Ser	Leu	Glu	Asn	His
			245						250					255	
Ala	His	Thr	Leu	Glu	Val	Glu	Leu	Glu	Ser	Leu	Lys	Val	Gln	Leu	Asp
			260					265						270	
Glu	Glu	Ser	Glu	Ala	Arg	Leu	Glu	Leu	Glu	Arg	Gln	Leu	Thr	Lys	Ala
		275					280					285			
Asn	Gly	Asp	Ala	Ala	Ser	Trp	Lys	Ser	Lys	Tyr	Glu	Ala	Glu	Leu	Gln
	290					295					300				
Ala	His	Ala	Asp	Glu	Val	Glu	Glu	Leu	Arg	Arg	Lys	Met	Ala	Gln	Lys
	305					310					315				320
Ile	Ser	Glu	Tyr	Glu	Glu	Gln	Leu	Glu	Ala	Leu	Leu	Asn	Lys	Cys	Ser
			325						330					335	
Ser	Leu	Glu	Lys	Gln	Lys	Ser	Arg	Leu	Gln	Ser	Glu	Val	Glu	Val	Leu
			340					345					350		
Ile	Met	Asp	Leu	Glu	Lys	Ala	Thr	Ala	His	Ala	Gln	Gln	Leu	Glu	Lys
		355					360						365		
Arg	Val	Ala	Gln	Leu	Glu	Lys	Ile	Asn	Leu	Asp	Leu	Lys	Asn	Lys	Leu
	370					375					380				
Glu	Glu	Val	Thr	Met	Leu	Met	Glu	Gln	Ala	Gln	Lys	Glu	Leu	Arg	Val
	385					390					395				400
Lys	Ile	Ala	Glu	Leu	Gln	Lys	Leu	Gln	His	Glu	Tyr	Glu	Lys	Leu	Arg
			405						410					415	
Asp	Gln	Arg	Asp	Gln	Leu	Ala	Arg	Glu	Asn	Lys	Lys	Leu	Thr	Asp	Asp
			420					425					430		
Leu	Ala	Glu	Ala	Lys	Ser	Gln	Leu	Asn	Asp	Ala	His	Arg	Arg	Ile	His
	435						440					445			
Glu	Gln	Glu	Ile	Glu	Ile	Lys	Arg	Leu	Glu	Asn	Glu	Arg	Asp	Glu	Leu
	450					455					460				
Ser	Ala	Ala	Tyr	Lys	Glu	Ala	Glu	Thr	Leu	Arg	Lys	Gln	Glu	Glu	Ala
	465					470					475				480
Lys	Asn	Gln	Arg	Leu	Ile	Ala	Glu	Leu	Ala	Gln	Val	Arg	His	Asp	Tyr
			485						490					495	
Glu	Lys	Arg	Leu	Ala	Gln	Lys	Asp	Glu	Glu	Ile	Glu	Ala	Leu	Arg	Lys
			500					505					510		
Gln	Tyr	Gln	Ile	Glu	Ile	Glu	Gln	Leu	Asn	Met	Arg	Leu	Ala	Glu	Ala
		515					520						525		

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Glu Ala Lys Leu Lys Thr Glu Ile Ala Arg Leu Lys Lys Lys Tyr Gln
 530 535 540
 Ala Gln Ile Thr Glu Leu Glu Leu Ser Leu Asp Ala Ala Asn Lys Ala
 545 550 555 560
 Asn Ile Asp Leu Gln Lys Thr Ile Lys Lys Gln Ala Leu Gln Ile Thr
 565 570 575
 Ser Glu Leu Gln Ala His Tyr Asp Glu Val His Arg Gln Leu Gln Gln
 580 585 590
 Ala Val Asp Gln Leu Gly Val Thr Gln Arg Arg Cys Gln Ala Leu Gln
 595 600 605
 Ala Glu Leu Glu Glu Met Arg Ile Ala Leu Glu Gln Ala Asn Arg Ala
 610 615 620
 Lys Arg Gln Ala Glu Gln Leu His Glu Glu Ala Val Val Arg Val Asn
 625 630 635 640
 Glu Leu Thr Thr Ile Asn Val Asn Leu Ala Ser Ala Lys Ser Lys Leu
 645 650 655
 Glu Ser Glu Phe Ser Ala Leu Gln Ala Asp Thr Met Lys Tyr Ile Lys
 660 665 670
 Asn Leu Glu Phe Leu Met Asn Glu Tyr Arg Asn Leu Gln Leu Asn Ser
 675 680 685
 Asn Leu Leu Lys Ile Cys
 690

<210> SEQ ID NO 27
 <211> LENGTH: 193
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 15_isol_TI

<400> SEQUENCE: 27

Met Lys Thr Ile Tyr Ala Ile Leu Ser Ile Met Ala Cys Ile Gly Leu
 1 5 10 15
 Met Asn Ala Ser Ile Lys Arg Asp His Asn Asp Tyr Ser Lys Asn Pro
 20 25 30
 Met Arg Ile Val Cys Tyr Val Gly Thr Trp Ser Val Tyr His Lys Val
 35 40 45
 Asp Pro Tyr Thr Ile Glu Asp Ile Asp Pro Phe Lys Cys Thr His Leu
 50 55 60
 Met Tyr Gly Phe Ala Lys Ile Asp Glu Tyr Lys Tyr Thr Ile Gln Val
 65 70 75 80
 Phe Asp Pro Tyr Gln Asp Asp Asn His Asn Ser Trp Glu Lys Arg Gly
 85 90 95
 Tyr Glu Arg Phe Asn Asn Leu Arg Leu Lys Asn Pro Glu Leu Thr Thr
 100 105 110
 Met Ile Ser Leu Gly Gly Trp Tyr Glu Gly Ser Glu Lys Tyr Ser Asp
 115 120 125
 Met Ala Ala Asn Pro Thr Tyr Arg Gln Gln Phe Ile Gln Ser Val Leu
 130 135 140
 Asp Phe Leu Gln Glu Tyr Lys Phe Asp Gly Leu Asp Trp Ile Gly Ser
 145 150 155 160
 Ile Leu Asp Leu Asp Trp Val Thr Arg Lys Ser Thr Asn Lys Thr Ile
 165 170 175
 Trp Leu Trp Leu Glu Asn Leu Lys Thr Leu Leu Asn Leu Met Ala Thr
 180 185 190

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Cys

<210> SEQ ID NO 28
 <211> LENGTH: 234
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 32_isol_TI

<400> SEQUENCE: 28

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Met Ser Thr Thr Asn Tyr Ser Val Asp His Arg Gly Ser Phe Asn Ser
1          5          10          15
Leu Asp Tyr Arg Ile Tyr Phe Lys Asp Asn Ser Asn Gly Lys Ile Ile
          20          25          30
Ser Pro Trp His Asp Ile Pro Leu Phe Val Asp Lys Ser Ala Lys His
          35          40          45
Tyr Asn Met Val Val Glu Ile Pro Arg Trp Thr Asn Glu Lys Met Glu
          50          55          60
Ile Ala Thr Ala Glu Pro Met Ser Pro Ile Lys Gln Asp Ile Lys Lys
65          70          75          80
Gly Ala Leu Arg Tyr Val Lys Asn Val Phe Pro His Lys Gly Tyr Ile
          85          90          95
Trp Asn Tyr Gly Ala Phe Pro Gln Thr Trp Glu Asn Pro Asn His Ile
          100          105          110
Asp Gln Asp Thr Lys Thr Lys Gly Asp Asn Asp Pro Ile Asp Val Ile
          115          120          125
Glu Ile Gly Ser Arg Val Ala Lys Arg Gly Asp Val Val Pro Val Lys
          130          135          140
Ile Leu Gly Thr Ile Ala Leu Ile Asp Glu Gly Glu Thr Asp Trp Lys
145          150          155          160
Ile Ile Ala Ile Asp Thr Arg Asp Glu Leu Ala Ser Gln Met Asn Asn
          165          170          175
Val Asp Asp Val Glu Lys Leu Leu Pro Gly Leu Leu Arg Ala Thr Val
          180          185          190
Glu Trp Phe Lys Ile Tyr Lys Ile Pro Asp Gly Lys Pro Ala Asn Lys
          195          200          205
Phe Ala Phe Asn Gly Glu Ala Lys Asp Arg Glu Leu Leu Lys Lys Ser
          210          215          220
Leu Lys Lys His Ile Asn Ile Gly Lys Lys
225          230

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<210> SEQ ID NO 29
 <211> LENGTH: 210
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 25_isol_TI

<400> SEQUENCE: 29

```

Met Val Arg Lys Phe Phe Val Gly Gly Asn Trp Lys Met Asn Gly Ser
1          5          10          15
Arg Ala Thr Asn Glu Asp Leu Ile Lys Thr Leu Ser Asn Gly Pro Leu
          20          25          30
Asp Pro Asn Thr Asp Val Val Val Gly Val Pro Ser Ile Tyr Met Ala
          35          40          45
Glu Val Arg Gln Lys Leu Pro Lys Thr Ile Gly Val Ala Ala Gln Asn
50          55          60

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Cys Tyr Lys Val Pro Lys Gly Ala Phe Thr Gly Glu Ile Ser Pro Ala
 65 70 75 80
 Met Ile Lys Asp Val Gly Ala Glu Trp Val Ile Leu Gly His Ser Glu
 85 90 95
 Arg Arg Asn Val Phe Gly Glu Ser Asp Gln Leu Ile Gly Glu Lys Val
 100 105 110
 Glu His Ala Leu Gln Glu Gly Leu His Val Ile Ala Cys Ile Gly Glu
 115 120 125
 Leu Leu Glu Glu Arg Glu Ala Gly Lys Thr Thr Glu Val Val Phe Arg
 130 135 140
 Gln Thr Gln Val Ile Ser Lys His Val Lys Asp Trp Ser Lys Val Val
 145 150 155 160
 Leu Ala Tyr Glu Pro Val Gly Pro Leu Val Leu Val Lys Gln Pro Val
 165 170 175
 His Asn Lys His Lys Lys Phe Ile Lys Asn Phe Asp Asn Gly Phe Leu
 180 185 190
 Lys Met Phe His His Lys Leu Pro Lys Gln Phe Glu Ser Phe Met Val
 195 200 205
 Val Gln
 210

<210> SEQ ID NO 30
 <211> LENGTH: 210
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 25_isol_TI

<400> SEQUENCE: 30

Met Val Arg Lys Phe Phe Val Gly Gly Asn Trp Lys Met Asn Gly Ser
 1 5 10 15
 Arg Ala Thr Asn Glu Asp Leu Ile Lys Thr Leu Ser Asn Gly Pro Leu
 20 25 30
 Asp Pro Asn Thr Asp Val Val Val Gly Val Pro Ser Ile Tyr Met Ala
 35 40 45
 Glu Val Arg Gln Lys Leu Pro Lys Thr Ile Gly Val Ala Ala Gln Asn
 50 55 60
 Cys Tyr Lys Val Pro Lys Gly Ala Phe Thr Gly Glu Ile Ser Pro Ala
 65 70 75 80
 Met Ile Lys Asp Val Gly Ala Glu Trp Val Ile Leu Gly His Ser Glu
 85 90 95
 Arg Arg Asn Val Phe Gly Glu Ser Asp Gln Leu Ile Gly Glu Lys Val
 100 105 110
 Glu His Ala Leu Gln Glu Gly Leu His Val Ile Ala Cys Ile Gly Glu
 115 120 125
 Leu Leu Glu Glu Arg Glu Ala Gly Lys Thr Thr Glu Val Val Phe Arg
 130 135 140
 Gln Thr Gln Val Ile Ser Lys His Val Lys Asp Trp Ser Lys Val Val
 145 150 155 160
 Leu Ala Tyr Glu Pro Val Trp Ala Ile Gly Thr Gly Lys Thr Ala Ser
 165 170 175
 Pro Gln Gln Ala Gln Glu Val His Gln Lys Leu Arg Gln Trp Phe Ser
 180 185 190
 Glu Met Phe His His Lys Leu Pro Lys Gln Phe Glu Ser Phe Met Val
 195 200 205

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Val Gln
210

<210> SEQ ID NO 31
<211> LENGTH: 460
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 16_iso1_TI

<400> SEQUENCE: 31

Met Ala Ala His Asp Lys Asn Phe Asp Val Ile Pro Ile Gly His Thr
1 5 10 15
Phe Phe Phe Ile Trp Arg Ile Lys Gln Phe Glu Leu Val Pro Val Pro
20 25 30
Lys Glu Asp Tyr Gly Lys Phe Tyr Lys Gly Asp Cys Tyr Ile Val Ala
35 40 45
Cys Cys Thr Glu Asn Pro Thr Gly Gly His Ser Lys Met Glu Ser Lys
50 55 60
Pro Ile Leu Asn Gly His Gly Tyr Cys His Ile His Phe Trp Ile Gly
65 70 75 80
Ser Glu Ser Thr Lys Asp Glu Ala Gly Val Ala Ala Ile Lys Ser Val
85 90 95
Glu Leu Asp Asp Phe Leu Gly Gly Tyr Pro Val Gln His Arg Glu Ile
100 105 110
Glu Glu Phe Glu Ser Arg Gln Phe Ser Ser Tyr Phe Lys Asn Gly Ile
115 120 125
Ile Tyr Leu Lys Gly Gly Tyr Glu Ser Gly Phe Thr Lys Met Ile Asp
130 135 140
Glu Leu Lys Pro Ser Leu Leu His Val Lys Gly Lys Lys Arg Pro Ile
145 150 155 160
Val Tyr Glu Cys Ala Glu Ile Ser Trp Lys Val Met Asn Asn Gly Asp
165 170 175
Val Phe Ile Leu Leu Val Pro Asn Phe Val Phe Val Trp Thr Gly Lys
180 185 190
His Ser Asn Arg Met Glu Arg Thr Thr Ala Ile Arg Val Ala Asn Asp
195 200 205
Leu Lys Ser Glu Leu Asn Arg Phe Lys Leu Ser Ser Val Ile Leu Glu
210 215 220
Asp Gly Lys Glu Val Glu Gln Thr Ser Gly Ala Glu Tyr Asp Ala Phe
225 230 235 240
Asn Lys Ala Leu Ser Leu Asp Lys Lys Asp Ile Asp Leu Lys Gln Met
245 250 255
Pro Lys Gly Tyr Asp Tyr Ala Ala Ser Asp Lys Ser Phe Glu Ser His
260 265 270
Glu Arg Ser Phe Val Thr Leu Tyr Lys Cys Phe Glu Gly Thr Glu Thr
275 280 285
Ile Asp Ile Ser Phe Val Lys Asn Gly Pro Leu Ser Arg Ala Asp Leu
290 295 300
Asp Thr Asn Asp Thr Phe Ile Val Glu Asn Gly Ser Glu Gly Leu Trp
305 310 315 320
Val Trp Val Gly Lys Lys Ala Thr Gln Lys Glu Arg Gln Ser Ala Ile
325 330 335
Lys Tyr Ala Met Glu Leu Ile Asn Lys Lys Lys Tyr Pro Asn Asn Thr
340 345 350

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Pro Val Thr Lys Val Leu Glu Gly Asp Glu Ser Val Glu Phe Lys Ser
 355 360 365

Leu Phe Glu Ser Trp Gln Met Ser Glu Gln Glu Lys Ile Thr Ser Ala
 370 375 380

Arg Leu Phe Arg Val Ser Arg Asn Gly Ile Phe Lys Gln Val Ala Asn
 385 390 395 400

Tyr Glu Pro Asp Asp Leu Glu Glu Asp Asn Ile Met Ile Leu Asp Val
 405 410 415

Met Asp Lys Ile Tyr Val Trp Ile Gly Asn Gln Phe Ala Glu Arg Ile
 420 425 430

Ala Asp Glu Ala His Val Asp Lys Val Ala Gln Arg Leu Tyr Lys Arg
 435 440 445

Ile Lys Val Ala Val Asn Phe Asp Gln Ile Arg Leu
 450 455 460

<210> SEQ ID NO 32
 <211> LENGTH: 60
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 26_isol_TI

<400> SEQUENCE: 32

Met Ala Leu Pro Arg Val Phe Phe Asp Ile Ala Ala Asp Asn Gln Pro
 1 5 10 15

Leu Gly Arg Ile Val Ile Glu Leu Arg Ser Asp Val Val Pro Lys Thr
 20 25 30

Ala Glu Ile Ser Val His Phe Ala Leu Val Lys Lys Asp Leu Val Leu
 35 40 45

Asn His Pro His Phe Ile Val Ser Tyr Pro Ile Leu
 50 55 60

<210> SEQ ID NO 33
 <211> LENGTH: 155
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 26_isol_TI

<400> SEQUENCE: 33

Met Ala Leu Pro Arg Val Phe Phe Asp Ile Ala Ala Asp Asn Gln Pro
 1 5 10 15

Leu Gly Arg Ile Val Ile Glu Leu Arg Ser Asp Val Val Pro Lys Thr
 20 25 30

Ala Glu Asn Phe Arg Ala Leu Cys Thr Gly Glu Lys Gly Phe Gly Phe
 35 40 45

Lys Ser Ser Ser Phe His Arg Ile Ile Pro Asn Phe Met Ile Gln Gly
 50 55 60

Gly Asp Phe Thr Asn His Asn Gly Thr Gly Gly Lys Ser Ile Tyr Gly
 65 70 75 80

Asn Lys Phe Ala Asp Glu Asn Phe Thr Leu Gln His Thr Gly Pro Gly
 85 90 95

Ile Met Ser Met Ala Asn Ala Gly Pro Asn Thr Asn Gly Ser Gln Phe
 100 105 110

Phe Ile Thr Thr Val Lys Thr Thr Trp Leu Asp Gly Lys His Val Val
 115 120 125

Phe Gly Ser Val Val Glu Gly Met Asp Ile Val Lys Arg Trp Lys Ala

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130	135	140
Met Ala His Asn Arg Val Asn His Pro Arg Lys		
145	150	155

<210> SEQ ID NO 34
 <211> LENGTH: 100
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 23_iso2_TI

<400> SEQUENCE: 34

Met Lys Phe Asn Ile Thr Ile Ala Phe Val Ser Leu Ala Ile Leu Ile		
1	5	10 15
His Ser Ser Tyr Ala Asp Ile Asp His Phe Asp Asn Asp Asp Gln Asn		
	20	25 30
Ser Ser Thr Ser Arg Pro Asp Asp Asp Pro Thr Thr Met Ile Asp Val		
	35	40 45
Gln Thr Thr Thr Val Gln Pro Ser Asp Glu Phe Glu Cys Pro Thr Arg		
	50	55 60
Phe Gly Tyr Phe Ala Asp Pro Lys Asp Pro Cys Lys Phe Ile Phe Val		
65	70	75 80
Gln Ile Gly Lys Leu Tyr Ile Lys Val Val Gln Val Ile Gln Asp Gly		
	85	90 95
Met Lys Lys Asn		
	100	

<210> SEQ ID NO 35
 <211> LENGTH: 101
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 13_isol_TI

<400> SEQUENCE: 35

Met Ala Ser Ile Glu Gly Lys Tyr Lys Leu Glu Lys Ser Glu Lys Phe		
1	5	10 15
Asp Glu Phe Leu Asp Lys Leu Gly Val Gly Phe Met Val Lys Thr Ala		
	20	25 30
Ala Lys Thr Leu Lys Pro Thr Phe Glu Val Ala Ile Glu Asn Asp Gln		
	35	40 45
Tyr Ile Phe Arg Ser Leu Ser Thr Phe Lys Asn Thr Glu Ala Lys Phe		
	50	55 60
Lys Leu Gly Glu Glu Phe Glu Glu Asp Arg Ala Asp Gly Lys Arg Val		
65	70	75 80
Lys Thr Val Ile Gln Lys Lys Val Thr Ile Asn Leu Phe Lys His Asn		
	85	90 95
Ser Val Ile Lys Lys		
	100	

<210> SEQ ID NO 36
 <211> LENGTH: 70
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 7_iso1_TI

<400> SEQUENCE: 36

Met Met Lys Phe Leu Leu Ile Ala Ala Val Ala Phe Val Ala Val Ser		
1	5	10 15

-continued

Ala Asp Pro Ile His Tyr Asp Lys Ile Thr Glu Glu Ile Asn Lys Ala
 20 25 30
 Ile Asp Asp Ala Ile Ala Ala Ile Glu Lys Ser Glu Thr Ile Asp Pro
 35 40 45
 Met Lys Val Pro Asp Leu Pro Ile Asn Ser Asn Val Met Leu Val Phe
 50 55 60
 Trp Ile Ser Lys Val Asn
 65 70

<210> SEQ ID NO 37
 <211> LENGTH: 335
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 27_isol_TI

<400> SEQUENCE: 37

Met Lys Phe Phe Leu Leu Ser Phe Val Leu Met Ile Val Ala Ala Thr
 1 5 10 15
 Ala Thr Tyr Ala Ala His Val Gly Ser Gly Ser Arg Asp Asn Asn Asn
 20 25 30
 Asn Lys Pro Val Pro Ala Glu Gly Phe Ala Lys Ala Ser Asn Glu Phe
 35 40 45
 Gly Phe His Leu Leu Lys Glu Val Ile Gln His Arg Ser Ser Ser Gly
 50 55 60
 Ser Arg Gly Ser Ser Glu Asn Val Leu Phe Ser Pro Tyr Ser Val Ala
 65 70 75 80
 Val Ala Leu Ser Met Val His Gln Gly Thr Gln Gly Ser Thr Ala Glu
 85 90 95
 Gln Phe Lys Arg Val Leu Tyr Tyr Asp Arg Val Gln Gln Leu Asn Gly
 100 105 110
 Gly Glu Tyr Gln Thr Val Ala Asn Ser Val Lys Gln Ile Gln Asn Gln
 115 120 125
 Ile Lys Gln Ser Asp Gln Ser Asn Gln Phe Asp Trp Gly Asn Met Leu
 130 135 140
 Met Val Asp Gln Gln Ile Pro Val Lys Asp Gln Tyr Lys Lys Ile Ile
 145 150 155 160
 Glu Gln Tyr Tyr Asp Gly Gln Val Met Ser Val Asp Phe Arg Lys Glu
 165 170 175
 Ser Lys Asn Val Met Glu Arg Ile Asn Gln Phe Val Ser Asn Lys Thr
 180 185 190
 His Gly Leu Ile Asp Arg Met Leu Glu Gln Pro Pro Ser Ala Asp Thr
 195 200 205
 Gly Leu Ala Leu Ile Asn Ala Val Tyr Phe Lys Gly Glu Trp Leu Lys
 210 215 220
 Pro Phe Asp Ser Met Arg Thr Glu Gln Ser Val Phe Tyr Gly His His
 225 230 235 240
 Gly Gln Glu Tyr Lys Asn Val Gln Tyr Ile Asn Gly Gln Gly Pro Tyr
 245 250 255
 Gly Tyr Val Glu Val Pro Gln Trp Asn Ser Asp Leu Ile Gln Leu Pro
 260 265 270
 Tyr Lys Gly Glu Asp Ile Ala Phe Tyr Gly Val Leu Pro Arg Glu Arg
 275 280 285
 Asn Met Ile Leu Thr Lys Phe Val Asn Gln Ser Ile Gln Leu Leu Leu
 290 295 300

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Met Lys Leu Leu Asp Lys Leu Leu Val Val Ser His Gln Leu Phe Ile
 305 310 315 320

Ser Arg Lys Leu Asn Ser Val His His Ile Asn Cys Arg Lys Phe
 325 330 335

<210> SEQ ID NO 38
 <211> LENGTH: 70
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 7_iso2_TI

<400> SEQUENCE: 38

Met Met Lys Phe Leu Leu Ile Ala Ala Val Ala Phe Val Ala Val Ser
 1 5 10 15
 Ala Asp Pro Ile His Tyr Asp Lys Ile Thr Glu Glu Ile Asn Lys Ala
 20 25 30
 Ile Asp Asp Ala Ile Ala Ala Ile Glu Lys Ser Glu Thr Ile Asp Pro
 35 40 45
 Met Lys Val Pro Asp Leu Pro Ile Asn Ser Asn Val Met Leu Val Phe
 50 55 60
 Trp Ile Ser Lys Val Asn
 65 70

<210> SEQ ID NO 39
 <211> LENGTH: 155
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 29_iso1_TI

<400> SEQUENCE: 39

Met Ala Leu Pro Arg Val Phe Phe Asp Ile Ala Ala Asp Asn Gln Pro
 1 5 10 15
 Leu Gly Arg Ile Val Ile Glu Leu Arg Ser Asp Val Val Pro Lys Thr
 20 25 30
 Ala Glu Asn Phe Arg Ala Leu Cys Thr Gly Glu Lys Gly Phe Gly Phe
 35 40 45
 Lys Ser Ser Ser Phe His Arg Ile Ile Pro Asn Phe Met Ile Gln Gly
 50 55 60
 Gly Asp Phe Thr Asn His Asn Gly Thr Gly Gly Lys Ser Ile Tyr Gly
 65 70 75 80
 Asn Lys Phe Ala Asp Glu Asn Phe Thr Leu Gln His Thr Gly Pro Gly
 85 90 95
 Ile Met Ser Met Ala Asn Ala Gly Pro Asn Thr Asn Gly Ser Gln Phe
 100 105 110
 Phe Ile Thr Thr Val Lys Thr Thr Trp Leu Asp Gly Lys His Val Val
 115 120 125
 Phe Gly Ser Val Val Glu Gly Met Asp Ile Val Lys Arg Trp Lys Ala
 130 135 140
 Met Ala His Asn Arg Val Asn His Pro Arg Lys
 145 150 155

<210> SEQ ID NO 40
 <211> LENGTH: 448
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 18_iso1_TI

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<400> SEQUENCE: 40

Met Thr Arg Phe Ser Leu Thr Val Leu Ala Val Leu Ala Ala Cys Phe
 1 5 10 15
 Gly Ser Asn Ile Arg Pro Asn Val Ala Thr Leu Glu Pro Lys Thr Val
 20 25 30
 Cys Tyr Tyr Glu Ser Trp Val His Trp Arg Gln Gly Glu Gly Lys Met
 35 40 45
 Asp Pro Glu Asp Ile Asp Thr Ser Leu Cys Thr His Ile Val Tyr Ser
 50 55 60
 Tyr Phe Gly Ile Asp Ala Ala Thr His Glu Ile Lys Leu Leu Asp Glu
 65 70 75 80
 Tyr Leu Met Lys Asp Leu His Asp Met Glu His Phe Thr Gln His Lys
 85 90 95
 Gly Asn Ala Lys Ala Met Ile Ala Val Gly Gly Ser Thr Met Ser Asp
 100 105 110
 Gln Phe Ser Lys Thr Ala Ala Val Glu His Tyr Arg Glu Thr Phe Val
 115 120 125
 Val Ser Thr Val Asp Leu Met Thr Arg Tyr Gly Phe Asp Gly Val Met
 130 135 140
 Ile Asp Trp Ser Gly Met Gln Ala Lys Asp Ser Asp Asn Phe Ile Lys
 145 150 155 160
 Leu Leu Asp Lys Phe Asp Glu Lys Phe Ala His Thr Ser Phe Val Met
 165 170 175
 Gly Val Thr Leu Pro Ala Thr Ile Ala Ser Tyr Asp Asn Tyr Asn Ile
 180 185 190
 Pro Ala Ile Ser Asn Tyr Val Asp Phe Met Asn Val Leu Ser Leu Asp
 195 200 205
 Tyr Thr Gly Ser Trp Ala His Thr Val Gly His Ala Ser Pro Phe Pro
 210 215 220
 Glu Gln Leu Lys Thr Leu Glu Ala Tyr His Lys Arg Gly Ala Pro Arg
 225 230 235 240
 His Lys Met Val Met Ala Val Pro Phe Tyr Ala Arg Thr Trp Ile Leu
 245 250 255
 Glu Lys Met Asn Lys Gln Asp Ile Gly Asp Lys Ala Ser Gly Pro Gly
 260 265 270
 Pro Arg Gly Gln Phe Thr Gln Thr Asp Gly Phe Leu Ser Tyr Asn Glu
 275 280 285
 Leu Cys Val Gln Ile Gln Ala Glu Thr Asn Ala Phe Thr Ile Thr Arg
 290 295 300
 Asp His Asp Asn Thr Ala Ile Tyr Ala Val Tyr Val His Ser Asn His
 305 310 315 320
 Ala Glu Trp Ile Ser Phe Glu Asp Arg His Thr Leu Gly Glu Lys Ala
 325 330 335
 Lys Asn Ile Thr Gln Gln Gly Tyr Ala Gly Met Ser Val Tyr Thr Leu
 340 345 350
 Ser Asn Glu Asp Val His Gly Val Cys Gly Asp Lys Asn Pro Leu Leu
 355 360 365
 His Ala Ile Gln Ser Asn Tyr Tyr His Gly Val Val Thr Glu Pro Thr
 370 375 380
 Val Val Thr Leu Pro Pro Val Thr His Thr Thr Glu His Val Thr Asp
 385 390 395 400
 Ile Pro Gly Val Phe His Cys His Glu Glu Gly Phe Phe Arg Asp Lys

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325					330					335					
Lys	Tyr	Ala	Met	Glu	Leu	Ile	Asn	Lys	Lys	Lys	Tyr	Pro	Asn	Asn	Thr
			340					345					350		
Pro	Val	Thr	Lys	Val	Leu	Glu	Gly	Asp	Glu	Ser	Val	Glu	Phe	Lys	Ser
			355					360					365		
Leu	Phe	Glu	Ser	Trp	Gln	Met	Ser	Glu	Gln	Glu	Lys	Ile	Thr	Ser	Ala
			370					375					380		
Arg	Leu	Phe	Arg	Val	Ser	Arg	Asn	Gly	Ile	Phe	Lys	Gln	Val	Ala	Asn
								390					395		
Tyr	Glu	Pro	Asp	Asp	Leu	Glu	Glu	Asp	Asn	Ile	Met	Ile	Leu	Asp	Val
								405					415		
Met	Asp	Lys	Ile	Tyr	Val	Trp	Ile	Gly	Asn	Gln	Phe	Ala	Glu	Arg	Ile
								420					430		
Ala	Asp	Glu	Ala	His	Val	Asp	Lys	Val	Ala	Gln	Arg	Leu	Tyr	Lys	Arg
								440					445		
Ile	Lys	Val	Ala	Val	Asn	Phe	Asp	Gln	Ile	Arg	Leu				
								450					460		

<210> SEQ ID NO 42
 <211> LENGTH: 70
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 7_iso3_TI

<400> SEQUENCE: 42

Met	Met	Lys	Phe	Leu	Leu	Ile	Ala	Ala	Val	Ala	Phe	Val	Ala	Val	Ser
1				5					10					15	
Ala	Asp	Pro	Ile	His	Tyr	Asp	Lys	Ile	Thr	Glu	Glu	Ile	Asn	Lys	Ala
				20					25					30	
Ile	Asp	Asp	Ala	Ile	Ala	Ala	Ile	Glu	Lys	Ser	Glu	Thr	Ile	Asp	Pro
				35					40					45	
Met	Lys	Val	Pro	Asp	Leu	Pro	Ile	Asn	Ser	Asn	Val	Met	Leu	Val	Phe
				50					55					60	
Trp	Ile	Ser	Lys	Val	Asn										
				65					70						

<210> SEQ ID NO 43
 <211> LENGTH: 139
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 20_isol_TI

<400> SEQUENCE: 43

Met	Val	Asp	Gln	Ala	Val	Ile	Asp	Lys	Leu	Glu	Ala	Gly	Phe	Gln	Lys
1				5					10					15	
Leu	Gln	Ser	Ser	Ala	Glu	Cys	His	Ser	Leu	Leu	Lys	Lys	Tyr	Leu	Thr
				20					25					30	
Arg	Asn	Val	Leu	Asp	Ala	Cys	Lys	Gly	Arg	Lys	Thr	Gly	Met	Gly	Ala
				35					40					45	
Thr	Leu	Val	Asp	Val	Val	Gln	Ser	Gly	Phe	Glu	Asn	Leu	Asp	Ser	Gly
				50					55					60	
Val	Gly	Leu	Tyr	Ala	Pro	Asp	Ala	Glu	Ser	Tyr	Thr	Leu	Phe	Lys	Glu
				65					70					75	
Leu	Phe	Asp	Pro	Val	Ile	Glu	Asp	Tyr	His	Lys	Gly	Phe	Lys	Pro	Thr
				85					90					95	

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Asp Lys His Pro Gln Thr Asp Phe Gly Asp Val Asn Thr Cys Val Met
 100 105 110

Trp Ile Gln Ile Met Asn Leu Ser Phe Gln His Val Tyr Val Val Ala
 115 120 125

Asp His Cys Lys Val Ile His Leu Ile His Ala
 130 135

<210> SEQ ID NO 44
 <211> LENGTH: 348
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 20_isol_TI

<400> SEQUENCE: 44

Met Val Asp Gln Ala Val Ile Asp Lys Leu Glu Ala Gly Phe Gln Lys
 1 5 10 15

Leu Gln Ser Ser Ala Glu Cys His Ser Leu Leu Lys Lys Tyr Leu Thr
 20 25 30

Arg Asn Val Leu Asp Ala Cys Lys Gly Arg Lys Thr Gly Met Gly Ala
 35 40 45

Thr Leu Val Asp Val Val Gln Ser Gly Phe Glu Asn Leu Asp Ser Gly
 50 55 60

Val Gly Leu Tyr Ala Pro Asp Ala Glu Ser Tyr Thr Leu Phe Lys Glu
 65 70 75 80

Leu Phe Asp Pro Val Ile Glu Asp Tyr His Lys Gly Phe Lys Pro Thr
 85 90 95

Asp Lys His Pro Gln Thr Asp Phe Gly Asp Val Asn Thr Leu Cys Asn
 100 105 110

Val Asp Pro Asn Asn Glu Phe Val Ile Ser Thr Arg Val Arg Cys Gly
 115 120 125

Arg Ser Leu Gln Gly Tyr Pro Phe Asn Pro Cys Leu Thr Glu Ala Gln
 130 135 140

Tyr Lys Glu Met Glu Glu Lys Val Lys Gly Gln Leu Asn Ser Phe Glu
 145 150 155 160

Gly Glu Leu Lys Gly Thr Tyr Tyr Pro Leu Leu Gly Met Asp Lys Ala
 165 170 175

Thr Gln Gln Gln Leu Ile Asp Asp His Phe Leu Phe Lys Glu Gly Asp
 180 185 190

Arg Phe Leu Gln Ala Ala Asn Ala Cys Arg Phe Trp Pro Val Gly Cys
 195 200 205

Gly Ile Phe His Asn Asp Asn Lys Thr Phe Leu Ile Trp Val Asn Glu
 210 215 220

Glu Asp His Leu Arg Ile Ile Ser Met Gln Lys Gly Gly Asp Leu Lys
 225 230 235 240

Gln Val Phe Ser Arg Leu Ile Asn Gly Val Asn His Ile Glu Lys Lys
 245 250 255

Leu Pro Phe Ser Arg Asp Asp Arg Leu Gly Phe Leu Thr Phe Cys Pro
 260 265 270

Thr Asn Leu Gly Thr Thr Ile Arg Ala Ser Val His Ile Lys Leu Pro
 275 280 285

Lys Leu Ala Ala Asp Arg Lys Asn Trp Lys Lys Leu Leu Ala Asn Ile
 290 295 300

Ile Tyr Lys Tyr Val Val Leu Pro Val Asn Thr Pro Lys Val Leu Ala
 305 310 315 320

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Arg Pro Ser Lys Arg Cys Lys Met Val Phe Leu Asn
340 345

<210> SEQ ID NO 46
 <211> LENGTH: 348
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 20_isol_TI

<400> SEQUENCE: 46

Met Val Asp Gln Ala Val Ile Asp Lys Leu Glu Ala Gly Phe Gln Lys
1 5 10 15
 Leu Gln Ser Ser Ala Glu Cys His Ser Leu Leu Lys Lys Tyr Leu Thr
20 25 30
 Arg Asn Val Leu Asp Ala Cys Lys Gly Arg Lys Thr Gly Met Gly Ala
35 40 45
 Thr Leu Val Asp Val Val Gln Ser Gly Phe Glu Asn Leu Asp Ser Gly
50 55 60
 Val Gly Leu Tyr Ala Pro Asp Ala Glu Ser Tyr Thr Leu Phe Lys Glu
65 70 75 80
 Leu Phe Asp Pro Val Ile Glu Asp Tyr His Lys Gly Phe Lys Pro Thr
85 90 95
 Asp Lys His Pro Gln Thr Asp Phe Gly Asp Val Asn Thr Leu Cys Asn
100 105 110
 Val Asp Pro Asn Asn Glu Phe Val Ile Ser Thr Arg Val Arg Cys Gly
115 120 125
 Arg Ser Leu Gln Gly Tyr Pro Phe Asn Pro Cys Leu Thr Glu Ala Gln
130 135 140
 Tyr Lys Glu Met Glu Glu Lys Val Lys Gly Gln Leu Asn Ser Phe Glu
145 150 155 160
 Gly Glu Leu Lys Gly Thr Tyr Tyr Pro Leu Leu Gly Met Asp Lys Ala
165 170 175
 Thr Gln Gln Gln Leu Ile Asp Asp His Phe Leu Phe Lys Glu Gly Asp
180 185 190
 Arg Phe Leu Gln Ala Ala Asn Ala Cys Arg Phe Trp Pro Val Gly Cys
195 200 205
 Gly Ile Phe His Asn Asp Asn Lys Thr Phe Leu Ile Trp Val Asn Glu
210 215 220
 Glu Asp His Leu Arg Ile Ile Ser Met Gln Lys Gly Gly Asp Leu Lys
225 230 235 240
 Gln Val Phe Ser Arg Leu Ile Asn Gly Val Asn His Ile Glu Lys Lys
245 250 255
 Leu Pro Phe Ser Arg Asp Asp Arg Leu Gly Phe Leu Thr Phe Cys Pro
260 265 270
 Thr Asn Leu Gly Thr Thr Ile Arg Ala Ser Val His Ile Lys Leu Pro
275 280 285
 Lys Leu Ala Ala Asp Arg Lys Lys Leu Glu Glu Val Ala Gly Lys Tyr
290 295 300
 Asn Leu Gln Val Arg Gly Thr Ala Gly Glu His Thr Glu Ser Val Gly
305 310 315 320
 Gly Val Thr Ile Ser Val Ile Asn Val Val Trp Val Leu Leu Asn Ile
325 330 335
 Arg Pro Ser Lys Arg Cys Lys Met Val Phe Leu Asn
340 345

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<210> SEQ ID NO 47
<211> LENGTH: 363
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 1_iso3_TI

<400> SEQUENCE: 47

Met Lys Phe Ala Leu Phe Val Val Ala Ser Leu Ile Ala Thr Val Tyr
 1          5          10          15

Gly Gln Ser His Gln Tyr Tyr His Thr Ser Gly Leu Arg Asn Leu Gly
 20          25          30

Gly Ser Tyr Tyr Arg Ser Ala Gly Ile Ser Gly Val Ala Gly Leu Gly
 35          40          45

Gly Leu Ala Tyr Gly Thr Gly Leu Gly Tyr Gly Thr Arg Tyr Gly Tyr
 50          55          60

Gly Ser Gly Leu Gly Tyr Gly Leu Gly Tyr Gly Leu Gly Tyr Gly Gln
 65          70          75          80

Ala Val Ala Leu Ala Pro Ala Gln Ala Val Gly Tyr Val Ala Ala Ala
 85          90          95

Pro Ala Val Ala Val Gln Ala Pro Ala Val Ser Tyr Ala Ala Ala Ala
 100         105         110

Pro Ala Val Gln Thr Val Ala Val Gln Ala Pro Ala Val Ser Tyr Ala
 115         120         125

Ala Ala Pro Ala Val Ala Val Gln Ala His Thr Ala Gln Val Ser Gly
 130         135         140

Pro Ile His Ala Ala Ile Glu Ser Arg Arg Thr Val Glu Val Ile Asp
 145         150         155         160

Gly Pro Ser Thr Gly Asp Ala Pro Val Ala Ser Thr Val Val Ile Gly
 165         170         175

Pro Asn Val Gln Pro Ile Asn Leu Glu Phe Gln Thr Gln Ala Ser Pro
 180         185         190

Leu Ala Ala Thr Gln Asn His Val Pro Thr Ala Pro Ala Glu Pro Gln
 195         200         205

Gln Ser Ser Tyr Glu Glu Gln Pro Asp Leu Leu Arg Gln Asp Ile Val
 210         215         220

Lys Pro Val Val Gln Asp Val His Glu Thr Ile Val Pro Phe Arg Arg
 225         230         235         240

Ile Thr Gln Glu Leu Lys Pro Val Gln Glu Ser Val His Gln Ile Leu
 245         250         255

Pro Arg Gly Gln Glu Arg Gly Phe Tyr Gln Gln Gln Gln Val Arg
 260         265         270

Val Ala Gln His Val Ala Ala Pro Ala Ala Val Ala Val Gln Pro Val
 275         280         285

Val Gln Ala Ala Pro Ala Ile Ser Ala Val Arg Val Ala Ala Ala Pro
 290         295         300

Ala Val Ala Tyr Ala Ala Pro Ala Val Ser Thr Val Ser Ala Ala Pro
 305         310         315         320

Ala Ala Ile Gly Val Ile Gly Val Gln Pro Ala Ala Gly Tyr Ile Gly
 325         330         335

Tyr Gly Ala Gly Tyr Gly Thr Gly Tyr Glu Gln Val Met Val Leu Leu
 340         345         350

Asn Thr Glu Pro Asp Met Val Ser Leu Ala Val
 355         360

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<210> SEQ ID NO 48
<211> LENGTH: 408
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 1_iso3_TI

<400> SEQUENCE: 48

Met Lys Phe Ala Leu Phe Val Val Ala Ser Leu Ile Ala Thr Val Tyr
 1          5          10          15

Gly Gln Ser His Gln Tyr Tyr His Thr Ser Gly Leu Arg Asn Leu Gly
 20          25          30

Gly Ser Tyr Tyr Arg Ser Ala Gly Ile Ser Gly Val Ala Gly Leu Gly
 35          40          45

Gly Leu Ala Tyr Gly Thr Gly Leu Gly Tyr Gly Thr Arg Tyr Gly Tyr
 50          55          60

Gly Ser Gly Leu Gly Tyr Gly Leu Gly Tyr Gly Leu Gly Tyr Gly Gln
 65          70          75          80

Ala Val Ala Leu Ala Pro Ala Gln Ala Val Gly Tyr Val Ala Ala Ala
 85          90          95

Pro Ala Val Ala Val Gln Ala Pro Ala Val Ser Tyr Ala Ala Ala Ala
 100         105         110

Pro Ala Val Gln Thr Val Ala Val Gln Ala Pro Ala Val Ser Tyr Ala
 115         120         125

Ala Ala Pro Ala Val Ala Val Gln Ala His Thr Ala Gln Val Ser Gly
 130         135         140

Pro Ile His Ala Ala Ile Glu Ser Arg Arg Thr Val Glu Val Ile Asp
 145         150         155         160

Gly Pro Ser Thr Gly Asp Ala Pro Val Ala Ser Thr Val Val Ile Gly
 165         170         175

Pro Asn Val Gln Pro Ile Asn Leu Glu Phe Gln Thr Gln Ala Ser Pro
 180         185         190

Leu Ala Ala Thr Gln Asn His Val Pro Thr Ala Pro Ala Glu Pro Gln
 195         200         205

Gln Ser Ser Tyr Glu Glu Gln Pro Asp Leu Leu Arg Gln Asp Ile Val
 210         215         220

Lys Pro Val Val Gln Asp Val His Glu Thr Ile Val Pro Phe Arg Arg
 225         230         235         240

Ile Thr Gln Glu Leu Lys Pro Val Gln Glu Ser Val His Gln Ile Leu
 245         250         255

Pro Arg Gly Gln Glu Arg Gly Phe Tyr Gln Gln Gln Gln Val Arg
 260         265         270

Val Ala Gln His Val Ala Ala Pro Ala Ala Val Ala Val Gln Pro Val
 275         280         285

Val Gln Ala Ala Pro Ala Ile Ser Ala Val Arg Val Ala Ala Ala Pro
 290         295         300

Ala Val Ala Tyr Ala Ala Pro Ala Val Ser Thr Val Ser Ala Ala Pro
 305         310         315         320

Ala Ala Ile Gly Val Ile Gly Val Gln Pro Ala Ala Gly Tyr Ile Gly
 325         330         335

Tyr Gly Ala Gly Tyr Gly Thr Gly Tyr Gly Thr Gly Tyr Gly Val Ala
 340         345         350

Lys Tyr Gly Thr Gly Tyr Gly Leu Thr Ser Gly Leu Ile Gly Val Ala
 355         360         365

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His Met Asp His His Ile Gln Tyr Asn Gln Pro Ala Thr Glu Leu Val
370 375 380

Met Val Thr Leu Pro Ile Ala Val Met Pro Thr Gln Ser Glu Lys Asn
385 390 395 400

Lys Leu Val Leu Pro Phe Ser Phe
405

<210> SEQ ID NO 49
<211> LENGTH: 193
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 15_iso2_TI

<400> SEQUENCE: 49

Met Lys Thr Ile Tyr Ala Ile Leu Ser Ile Met Ala Cys Ile Gly Leu
1 5 10 15

Met Asn Ala Ser Ile Lys Arg Asp His Asn Asp Tyr Ser Lys Asn Pro
20 25 30

Met Arg Ile Val Cys Tyr Val Gly Thr Trp Ser Val Tyr His Lys Val
35 40 45

Asp Pro Tyr Thr Ile Glu Asp Ile Asp Pro Phe Lys Cys Thr His Leu
50 55 60

Met Tyr Gly Phe Ala Lys Ile Asp Glu Tyr Lys Tyr Thr Ile Gln Val
65 70 75 80

Phe Asp Pro Tyr Gln Asp Asp Asn His Asn Ser Trp Glu Lys Arg Gly
85 90 95

Tyr Glu Arg Phe Asn Asn Leu Arg Leu Lys Asn Pro Glu Leu Thr Thr
100 105 110

Met Ile Ser Leu Gly Gly Trp Tyr Glu Gly Ser Glu Lys Tyr Ser Asp
115 120 125

Met Ala Ala Asn Pro Thr Tyr Arg Gln Gln Phe Ile Gln Ser Val Leu
130 135 140

Asp Phe Leu Gln Glu Tyr Lys Phe Asp Gly Leu Asp Trp Ile Gly Ser
145 150 155 160

Ile Leu Asp Leu Asp Trp Val Thr Arg Lys Ser Thr Asn Lys Thr Ile
165 170 175

Trp Leu Trp Leu Glu Asn Leu Lys Thr Leu Leu Asn Leu Met Ala Thr
180 185 190

Cys

<210> SEQ ID NO 50
<211> LENGTH: 348
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 20_iso2_TI

<400> SEQUENCE: 50

Met Val Asp Gln Ala Thr Leu Ser Lys Leu Glu Ala Gly Phe Gln Lys
1 5 10 15

Leu Gln Asn Ala Gln Asp Cys His Ser Leu Leu Lys Lys Tyr Leu Thr
20 25 30

Arg Asp Val Leu Asp Gln Leu Lys Thr Lys Lys Thr Asp Met Gly Ala
35 40 45

Thr Leu Leu Asp Val Ile Gln Ser Gly Val Glu Asn Leu Asp Ser Gly
50 55 60

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Val Gly Ile Tyr Ala Pro Asp Ala Gln Ser Tyr Lys Thr Phe Ala Ala
 65 70 75 80
 Leu Phe Asp Pro Ile Ile Asp Asp Tyr His Lys Gly Phe Lys Pro Thr
 85 90 95
 Asp Lys His Pro Gln Thr Asp Phe Gly Asn Ile Glu His Phe Val Asn
 100 105 110
 Val Asp Pro Lys Asn Glu Tyr Val Ile Ser Thr Arg Val Arg Cys Gly
 115 120 125
 Arg Ser Leu Lys Gly Tyr Pro Phe Asn Pro Met Leu Thr Glu Ala Gln
 130 135 140
 Tyr Lys Glu Met Glu Thr Lys Val Lys Gly Gln Leu Ala Thr Phe Glu
 145 150 155 160
 Gly Glu Leu Lys Gly Thr Tyr Tyr Pro Leu Leu Gly Met Asp Lys Ala
 165 170 175
 Thr Gln Gln Lys Leu Ile Asp Asp His Phe Leu Phe Lys Glu Gly Asp
 180 185 190
 Arg Phe Leu Gln Ala Ala Asn Ala Cys Arg Tyr Trp Pro Val Gly Arg
 195 200 205
 Gly Ile Phe His Asn Asp Lys Lys Thr Phe Leu Met Trp Val Asn Glu
 210 215 220
 Glu Asp His Leu Arg Ile Ile Ser Met Gln Lys Gly Gly Asp Leu Lys
 225 230 235 240
 Glu Val Phe Gly Arg Leu Val Lys Ala Val Lys His Ile Glu Gln Lys
 245 250 255
 Ile Pro Phe Ser Arg Asp Asp Arg Leu Gly Tyr Leu Thr Phe Cys Pro
 260 265 270
 Thr Asn Leu Gly Thr Thr Ile Arg Ala Ser Val His Ile Lys Leu Pro
 275 280 285
 Lys Leu Ala Ala Asp Arg Lys Lys Leu Glu Glu Val Ala Ala Arg Tyr
 290 295 300
 Asn Leu Gln Val Arg Gly Thr Ala Gly Glu His Thr Glu Ser Val Val
 305 310 315 320
 Val Ser Met Ile Leu Val Thr Asn Asp Glu Trp Val Ser Pro Asn Thr
 325 330 335
 Lys Leu Leu Arg Lys Cys Lys Met Ala Ser Leu Asn
 340 345

<210> SEQ ID NO 51

<211> LENGTH: 348

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Der f 20_iso2_TI

<400> SEQUENCE: 51

Met Val Asp Gln Ala Thr Leu Ser Lys Leu Glu Ala Gly Phe Gln Lys
 1 5 10 15
 Leu Gln Asn Ala Gln Asp Cys His Ser Leu Leu Lys Lys Tyr Leu Thr
 20 25 30
 Arg Asp Val Leu Asp Gln Leu Lys Thr Lys Lys Thr Asp Met Gly Ala
 35 40 45
 Thr Leu Leu Asp Val Ile Gln Ser Gly Val Glu Asn Leu Asp Ser Gly
 50 55 60
 Val Gly Ile Tyr Ala Pro Asp Ala Gln Ser Tyr Lys Thr Phe Ala Ala
 65 70 75 80

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Ala Glu Thr Ser Ala Cys Arg Ile Asn Ser Val Asn Val Pro Ser Glu
100 105 110

Leu Asp Leu Arg Ser Leu Arg Thr Val Thr Pro Ile Arg Met Gln Gly
115 120 125

Gly Cys Gly Ser Cys Trp Ala Phe Ser Gly Val Ala Ala Thr Glu Ser
130 135 140

Ala Tyr Leu Ala Tyr Arg Asn Thr Ser Leu Asp Leu Ser Glu Gln Glu
145 150 155 160

Leu Val Asp Cys Ala Ser Gln His Gly Cys His Gly Asp Thr Ile Pro
165 170 175

Arg Gly Ile Glu Tyr Ile Gln Gln Asn Gly Val Val Glu Glu Arg Ser
180 185 190

Tyr Pro Tyr Val Ala Arg Glu Gln Gln Cys Arg Arg Pro Asn Ser Gln
195 200 205

His Tyr Gly Ile Ser Asn Tyr Cys Gln Ile Tyr Pro Pro Asp Val Lys
210 215 220

Gln Ile Arg Glu Ala Leu Thr Gln Thr His Thr Ala Ile Ala Val Ile
225 230 235 240

Ile Gly Ile Lys Asp Leu Arg Ala Phe Gln His Tyr Asp Gly Arg Thr
245 250 255

Ile Ile Gln His Asp Asn Gly Tyr Gln Pro Asn Tyr His Ala Val Asn
260 265 270

Ile Val Gly Tyr Gly Ser Thr Gln Gly Val Asp Tyr Trp Ile Val Arg
275 280 285

Asn Ser Trp Asp Thr Thr Trp Gly Asp Ser Gly Tyr Gly Tyr Phe Gln
290 295 300

Ala Gly Asn Asn Leu Met Met Ile Glu Gln Tyr Pro Tyr Val Val Ile
305 310 315 320

Met

<210> SEQ ID NO 53
<211> LENGTH: 321
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 1_iso2

<400> SEQUENCE: 53

Met Lys Phe Val Leu Ala Ile Ala Ser Leu Leu Val Leu Ser Thr Val
1 5 10 15

Tyr Ala Arg Pro Ala Ser Ile Lys Thr Phe Glu Glu Phe Lys Lys Ala
20 25 30

Phe Asn Lys Asn Tyr Ala Thr Val Glu Glu Glu Glu Val Ala Arg Lys
35 40 45

Asn Phe Leu Glu Ser Leu Lys Tyr Val Glu Ala Asn Lys Gly Ala Ile
50 55 60

Asn His Leu Ser Asp Leu Ser Leu Asp Glu Phe Lys Asn Arg Tyr Leu
65 70 75 80

Met Ser Ala Glu Ala Phe Glu Gln Leu Lys Thr Gln Phe Asp Leu Asn
85 90 95

Ala Glu Thr Ser Ala Cys Arg Ile Asn Ser Val Asn Val Pro Ser Glu
100 105 110

Leu Asp Leu Arg Ser Leu Arg Thr Val Thr Pro Ile Arg Met Gln Gly
115 120 125

Gly Cys Gly Ser Cys Trp Ala Phe Ser Gly Val Ala Ala Thr Glu Ser

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Glu Leu Thr Cys Thr
180

<210> SEQ ID NO 55
<211> LENGTH: 876
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 11_isol

<400> SEQUENCE: 55

Met Ser Ala Arg Thr Ala Lys Tyr Met Tyr Arg Ser Ser Gly Ala Gly
1 5 10 15
Ala Ser Gly Asp Ile Ser Val Glu Tyr Gly Thr Asp Leu Gly Ala Leu
20 25 30
Thr Arg Leu Glu Asp Lys Ile Arg Leu Leu Ser Asp Asp Leu Glu Ser
35 40 45
Glu Arg Glu Met Arg Gln Arg Ile Glu Arg Glu Lys Ala Glu Leu Gln
50 55 60
Ile Gln Val Met Ser Leu Gly Glu Arg Leu Glu Glu Ala Glu Gly Ser
65 70 75 80
Ser Glu Ser Val Thr Glu Met Asn Lys Lys Arg Asp Ser Glu Leu Ala
85 90 95
Lys Leu Arg Lys Leu Leu Glu Asp Val His Ile Glu Ser Glu Glu Thr
100 105 110
Ala His His Leu Arg Gln Lys His Gln Ala Ala Ile Gln Glu Met Gln
115 120 125
Asp Gln Leu Asp Gln Leu Gln Lys Ala Lys Asn Lys Ser Asp Lys Glu
130 135 140
Lys Gln Lys Phe Gln Ala Glu Val Phe Glu Leu Leu Ala Gln Leu Glu
145 150 155 160
Thr Ala Asn Lys Glu Lys Leu Thr Ala Leu Lys Asn Val Glu Lys Leu
165 170 175
Glu Tyr Thr Val His Glu Leu Asn Ile Lys Ile Glu Glu Ile Asn Arg
180 185 190
Thr Val Ile Glu Leu Thr Ser His Lys Gln Arg Leu Ser Gln Glu Asn
195 200 205
Thr Glu Leu Ile Lys Glu Val His Glu Val Lys Leu Gln Leu Asp Asn
210 215 220
Ala Asn His Leu Lys Thr Gln Ile Ala Gln Gln Leu Glu Asp Thr Arg
225 230 235 240
His Arg Leu Glu Glu Glu Arg Lys Arg Ala Ser Leu Glu Asn His
245 250 255
Ala His Thr Leu Glu Val Glu Leu Glu Ser Leu Lys Val Gln Leu Asp
260 265 270
Glu Glu Ser Glu Ala Arg Leu Glu Leu Glu Arg Gln Leu Thr Lys Ala
275 280 285
Asn Gly Asp Ala Ala Ser Trp Lys Ser Lys Tyr Glu Ala Glu Leu Gln
290 295 300
Ala His Ala Asp Glu Val Glu Glu Leu Arg Arg Lys Met Ala Gln Lys
305 310 315 320
Ile Ser Glu Tyr Glu Glu Gln Leu Glu Ala Leu Leu Asn Lys Cys Ser
325 330 335
Ser Leu Glu Lys Gln Lys Ser Arg Leu Gln Ser Glu Val Glu Val Leu
340 345 350

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Ile Met Asp Leu Glu Lys Ala Thr Ala His Ala Gln Gln Leu Glu Lys
 355 360 365

Arg Val Ala Gln Leu Glu Lys Ile Asn Leu Asp Leu Lys Asn Lys Leu
 370 375 380

Glu Glu Val Thr Met Leu Met Glu Gln Ala Gln Lys Glu Leu Arg Val
 385 390 395 400

Lys Ile Ala Glu Leu Gln Lys Leu Gln His Glu Tyr Glu Lys Leu Arg
 405 410 415

Asp Gln Arg Asp Gln Leu Ala Arg Glu Asn Lys Lys Leu Thr Asp Asp
 420 425 430

Leu Ala Glu Ala Lys Ser Gln Leu Asn Asp Ala His Arg Arg Ile His
 435 440 445

Glu Gln Glu Ile Glu Ile Lys Arg Leu Glu Asn Glu Arg Asp Glu Leu
 450 455 460

Ser Ala Ala Tyr Lys Glu Ala Glu Thr Leu Arg Lys Gln Glu Glu Ala
 465 470 475 480

Lys Asn Gln Arg Leu Ile Ala Glu Leu Ala Gln Val Arg His Asp Tyr
 485 490 495

Glu Lys Arg Leu Ala Gln Lys Asp Glu Glu Ile Glu Ala Leu Arg Lys
 500 505 510

Gln Tyr Gln Ile Glu Ile Glu Gln Leu Asn Met Arg Leu Ala Glu Ala
 515 520 525

Glu Ala Lys Leu Lys Thr Glu Ile Ala Arg Leu Lys Lys Lys Tyr Gln
 530 535 540

Ala Gln Ile Thr Glu Leu Glu Leu Ser Leu Asp Ala Ala Asn Lys Ala
 545 550 555 560

Asn Ile Asp Leu Gln Lys Thr Ile Lys Lys Gln Ala Leu Gln Ile Thr
 565 570 575

Ser Glu Leu Gln Ala His Tyr Asp Glu Val His Arg Gln Leu Gln Gln
 580 585 590

Ala Val Asp Gln Leu Gly Val Thr Gln Arg Arg Cys Gln Ala Leu Gln
 595 600 605

Ala Glu Leu Glu Glu Met Arg Ile Ala Leu Glu Gln Ala Asn Arg Ala
 610 615 620

Lys Arg Gln Ala Glu Gln Leu His Glu Glu Ala Val Val Arg Val Asn
 625 630 635 640

Glu Leu Thr Thr Ile Asn Val Asn Leu Ala Ser Ala Lys Ser Lys Leu
 645 650 655

Glu Ser Glu Phe Ser Ala Leu Gln Ala Asp Tyr Asp Glu Val His Lys
 660 665 670

Glu Leu Arg Ile Ser Asp Glu Arg Val Gln Lys Leu Thr Ile Glu Leu
 675 680 685

Lys Ser Thr Lys Asp Leu Leu Ile Glu Glu Gln Glu Arg Leu Val Lys
 690 695 700

Leu Glu Thr Val Lys Lys Ser Leu Glu Gln Glu Val Arg Thr Leu His
 705 710 715 720

Val Arg Ile Glu Glu Val Glu Ala Asn Ala Leu Ala Gly Gly Lys Arg
 725 730 735

Val Ile Ala Lys Leu Glu Ser Arg Ile Arg Asp Val Glu Ile Glu Val
 740 745 750

Glu Glu Glu Arg Arg Arg His Ala Glu Thr Asp Lys Met Leu Arg Lys
 755 760 765

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Lys Asp His Arg Val Lys Glu Leu Leu Gln Asn Glu Glu Asp His
770 775 780

Lys Gln Ile Gln Leu Leu Gln Glu Met Thr Asp Lys Leu Asn Glu Lys
785 790 795 800

Val Lys Val Tyr Lys Arg Gln Met Gln Glu Gln Glu Gly Met Ser Gln
805 810 815

Gln Asn Leu Thr Arg Val Arg Arg Phe Gln Arg Glu Leu Glu Ala Ala
820 825 830

Glu Asp Arg Ala Asp Gln Ala Glu Ser Asn Leu Ser Phe Ile Arg Ala
835 840 845

Lys His Arg Ser Trp Val Thr Thr Ser Gln Val Pro Gly Gly Thr Arg
850 855 860

Gln Val Phe Thr Thr Gln Glu Glu Thr Thr Asn Tyr
865 870 875

<210> SEQ ID NO 56
 <211> LENGTH: 556
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 15_isol

<400> SEQUENCE: 56

Met Lys Thr Ile Tyr Ala Ile Leu Ser Ile Met Ala Cys Ile Gly Leu
1 5 10 15

Met Asn Ala Ser Ile Lys Arg Asp His Asn Asp Tyr Ser Lys Asn Pro
20 25 30

Met Arg Ile Val Cys Tyr Val Gly Thr Trp Ser Val Tyr His Lys Val
35 40 45

Asp Pro Tyr Thr Ile Glu Asp Ile Asp Pro Phe Lys Cys Thr His Leu
50 55 60

Met Tyr Gly Phe Ala Lys Ile Asp Glu Tyr Lys Tyr Thr Ile Gln Val
65 70 75 80

Phe Asp Pro Tyr Gln Asp Asp Asn His Asn Ser Trp Glu Lys Arg Gly
85 90 95

Tyr Glu Arg Phe Asn Asn Leu Arg Leu Lys Asn Pro Glu Leu Thr Thr
100 105 110

Met Ile Ser Leu Gly Gly Trp Tyr Glu Gly Ser Glu Lys Tyr Ser Asp
115 120 125

Met Ala Ala Asn Pro Thr Tyr Arg Gln Gln Phe Ile Gln Ser Val Leu
130 135 140

Asp Phe Leu Gln Glu Tyr Lys Phe Asp Gly Leu Asp Leu Asp Trp Glu
145 150 155 160

Tyr Pro Gly Ser Arg Leu Gly Asn Pro Lys Ile Asp Lys Gln Asn Tyr
165 170 175

Leu Ala Leu Val Arg Glu Leu Lys Asp Ala Phe Glu Pro His Gly Tyr
180 185 190

Leu Leu Thr Ala Ala Val Ser Pro Gly Lys Asp Lys Ile Asp Arg Ala
195 200 205

Tyr Asp Ile Lys Glu Leu Asn Lys Leu Phe Asp Trp Met Asn Val Met
210 215 220

Thr Tyr Asp Tyr His Gly Gly Trp Glu Asn Phe Tyr Gly His Asn Ala
225 230 235 240

Pro Leu Tyr Lys Arg Pro Asp Glu Thr Asp Glu Leu His Thr Tyr Phe
245 250 255

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Asn Val Asn Tyr Thr Met His Tyr Tyr Leu Asn Asn Gly Ala Thr Arg
 260 265 270

Asp Lys Leu Val Met Gly Val Pro Phe Tyr Gly Arg Ala Trp Ser Ile
 275 280 285

Glu Asp Arg Ser Lys Leu Lys Leu Gly Asp Pro Ala Lys Gly Met Ser
 290 295 300

Pro Pro Gly Phe Ile Ser Gly Glu Glu Gly Val Leu Ser Tyr Ile Glu
 305 310 315 320

Leu Cys Gln Leu Phe Gln Lys Glu Glu Trp His Ile Gln Tyr Asp Glu
 325 330 335

Tyr Tyr Asn Ala Pro Tyr Gly Tyr Asn Asp Lys Ile Trp Val Gly Tyr
 340 345 350

Asp Asp Leu Ala Ser Ile Ser Cys Lys Leu Ala Phe Leu Lys Glu Leu
 355 360 365

Gly Val Ser Gly Val Ile Val Trp Ser Leu Glu Asn Asp Asp Phe Lys
 370 375 380

Gly His Cys Gly Pro Lys Asn Pro Leu Leu Asn Lys Val His Asn Met
 385 390 400

Ile Asn Gly Asp Glu Lys Asn Ser Phe Glu Cys Ile Leu Gly Pro Ser
 405 410 415

Thr Thr Thr Pro Thr Pro Thr Thr Thr Pro Thr Thr Thr Pro Thr Pro
 420 425 430

Ser Pro Thr Thr Pro Thr Pro Ser Pro Thr Thr Pro Thr Thr Thr Pro
 435 440 445

Ser Pro Thr Thr Pro Thr Pro Ser Pro Thr Thr Pro Thr Thr Thr Pro
 450 455 460

Ser Pro Thr Thr Pro Thr Pro Thr Thr Pro Thr Pro Thr Pro Thr Thr
 465 470 475 480

Ser Thr Pro Ser Pro Thr Thr Thr Glu His Thr Ser Glu Thr Pro Lys
 485 490 495

Tyr Thr Thr Tyr Val Asp Gly His Leu Ile Lys Cys Tyr Lys Glu Gly
 500 505 510

Asp Ile Pro His Pro Thr Asn Ile His Lys Tyr Leu Val Cys Glu Phe
 515 520 525

Val Asn Gly Gly Trp Trp Val His Ile Met Pro Cys Pro Pro Gly Thr
 530 535 540

Ile Trp Cys Gln Glu Lys Leu Thr Cys Ile Gly Glu
 545 550 555

<210> SEQ ID NO 57
 <211> LENGTH: 296
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 32_isol

<400> SEQUENCE: 57

Met Ser Thr Thr Asn Tyr Ser Val Asp His Arg Gly Ser Phe Asn Ser
 1 5 10 15

Leu Asp Tyr Arg Ile Tyr Phe Lys Asp Asn Ser Asn Gly Lys Ile Ile
 20 25 30

Ser Pro Trp His Asp Ile Pro Leu Phe Val Asp Lys Ser Ala Lys His
 35 40 45

Tyr Asn Met Val Val Glu Ile Pro Arg Trp Thr Asn Glu Lys Met Glu
 50 55 60

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Leu Leu Glu Glu Arg Glu Ala Gly Lys Thr Thr Glu Val Val Phe Arg
 130 135 140
 Gln Thr Gln Val Ile Ser Lys His Val Lys Asp Trp Ser Lys Val Val
 145 150 155 160
 Leu Ala Tyr Glu Pro Val Trp Ala Ile Gly Thr Gly Lys Thr Ala Ser
 165 170 175
 Pro Gln Gln Ala Gln Glu Val His Gln Lys Leu Arg Gln Trp Phe Ser
 180 185 190
 Glu Asn Val Ser Pro Gln Ile Ala Glu Thr Ile Arg Ile Ile Tyr Gly
 195 200 205
 Gly Ser Val Thr Ala Asn Asn Ala Lys Glu Leu Ala Ser Gln Ala Asp
 210 215 220
 Val Asp Gly Phe Leu Val Gly Gly Ala Ser Leu Lys Pro Glu Phe Val
 225 230 235 240
 Gln Ile Val Asn Ala Arg Gln
 245

<210> SEQ ID NO 59
 <211> LENGTH: 480
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 16_isol

<400> SEQUENCE: 59

Met Ala Ala His Asp Lys Asn Phe Asp Val Ile Pro Ile Gly His Thr
 1 5 10 15
 Phe Phe Phe Ile Trp Arg Ile Lys Gln Phe Glu Leu Val Pro Val Pro
 20 25 30
 Lys Glu Asp Tyr Gly Lys Phe Tyr Lys Gly Asp Cys Tyr Ile Val Ala
 35 40 45
 Cys Cys Thr Glu Asn Pro Thr Gly Gly His Ser Lys Met Glu Ser Lys
 50 55 60
 Pro Ile Leu Asn Gly His Gly Tyr Cys His Ile His Phe Trp Ile Gly
 65 70 75 80
 Ser Glu Ser Thr Lys Asp Glu Ala Gly Val Ala Ala Ile Lys Ser Val
 85 90 95
 Glu Leu Asp Asp Phe Leu Gly Gly Tyr Pro Val Gln His Arg Glu Ile
 100 105 110
 Glu Glu Phe Glu Ser Arg Gln Phe Ser Ser Tyr Phe Lys Asn Gly Ile
 115 120 125
 Ile Tyr Leu Lys Gly Gly Tyr Glu Ser Gly Phe Thr Lys Met Ile Asp
 130 135 140
 Glu Leu Lys Pro Ser Leu Leu His Val Lys Gly Lys Lys Arg Pro Ile
 145 150 155 160
 Val Tyr Glu Cys Ala Glu Ile Ser Trp Lys Val Met Asn Asn Gly Asp
 165 170 175
 Val Phe Ile Leu Leu Val Pro Asn Phe Val Phe Val Trp Thr Gly Lys
 180 185 190
 His Ser Asn Arg Met Glu Arg Thr Thr Ala Ile Arg Val Ala Asn Asp
 195 200 205
 Leu Lys Ser Glu Leu Asn Arg Phe Lys Leu Ser Ser Val Ile Leu Glu
 210 215 220
 Asp Gly Lys Glu Val Glu Gln Thr Ser Gly Ala Glu Tyr Asp Ala Phe
 225 230 235 240

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Asn Lys Ala Leu Ser Leu Asp Lys Lys Asp Ile Asp Leu Lys Gln Met
      245                               250                255
Pro Lys Gly Tyr Asp Tyr Ala Ala Ser Asp Lys Ser Phe Glu Ser His
      260                               265                270
Glu Arg Ser Phe Val Thr Leu Tyr Lys Cys Phe Glu Gly Thr Glu Thr
      275                               280                285
Ile Asp Ile Ser Phe Val Lys Asn Gly Pro Leu Ser Arg Ala Asp Leu
      290                               295                300
Asp Thr Asn Asp Thr Phe Ile Val Glu Asn Gly Ser Glu Gly Leu Trp
      305                               310                315
Val Trp Val Gly Lys Lys Ala Thr Gln Lys Glu Arg Gln Ser Ala Ile
      325                               330                335
Lys Tyr Ala Met Glu Leu Ile Asn Lys Lys Lys Tyr Pro Asn Asn Thr
      340                               345                350
Pro Val Thr Lys Val Leu Glu Gly Asp Glu Ser Val Glu Phe Lys Ser
      355                               360                365
Leu Phe Glu Ser Trp Gln Met Ser Glu Gln Glu Lys Ile Thr Ser Ala
      370                               375                380
Arg Leu Phe Arg Val Ser Arg Asn Gly Ile Phe Lys Gln Val Ala Asn
      385                               390                395
Tyr Glu Pro Asp Asp Leu Glu Glu Asp Asn Ile Met Ile Leu Asp Val
      405                               410                415
Met Asp Lys Ile Tyr Val Trp Ile Gly Asn Gln Phe Ala Glu Arg Ile
      420                               425                430
Ala Asp Glu Ala His Val Asp Lys Val Ala Gln Arg Phe Ile Gln Glu
      435                               440                445
Asp Lys Ser Gly Arg Lys Phe Arg Pro Asn Gln Ile Ile Lys Leu Lys
      450                               455                460
Gln Gly Ser Glu Asp Gly Ala Phe Lys Ser Tyr Phe Pro Lys Trp Asn
      465                               470                475

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<210> SEQ ID NO 60
<211> LENGTH: 164
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 26_isol

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<400> SEQUENCE: 60

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Met Ala Leu Pro Arg Val Phe Phe Asp Ile Ala Ala Asp Asn Gln Pro
 1      5      10      15
Leu Gly Arg Ile Val Ile Glu Leu Arg Ser Asp Val Val Pro Lys Thr
      20      25      30
Ala Glu Asn Phe Arg Ala Leu Cys Thr Gly Glu Lys Gly Phe Gly Phe
      35      40      45
Lys Ser Ser Ser Phe His Arg Ile Ile Pro Asn Phe Met Ile Gln Gly
      50      55      60
Gly Asp Phe Thr Asn His Asn Gly Thr Gly Gly Lys Ser Ile Tyr Gly
      65      70      75      80
Asn Lys Phe Ala Asp Glu Asn Phe Thr Leu Gln His Thr Gly Pro Gly
      85      90      95
Ile Met Ser Met Ala Asn Ala Gly Pro Asn Thr Asn Gly Ser Gln Phe
      100     105     110
Phe Ile Thr Thr Val Lys Thr Thr Trp Leu Asp Gly Lys His Val Val
      115     120     125

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Phe Gly Ser Val Val Glu Gly Met Asp Ile Val Lys Lys Val Glu Ser
130 135 140

Tyr Gly Ser Gln Ser Gly Lys Pro Ser Lys Lys Val Thr Ile Ala Asn
145 150 155 160

Cys Gly Gln Leu

<210> SEQ ID NO 61
<211> LENGTH: 104
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 23_iso2

<400> SEQUENCE: 61

Met Lys Phe Asn Ile Thr Ile Ala Phe Val Ser Leu Ala Ile Leu Ile
1 5 10 15

His Ser Ser Tyr Ala Asp Ile Asp His Phe Asp Asn Asp Asp Gln Asn
20 25 30

Ser Ser Thr Ser Arg Pro Asp Asp Asp Pro Thr Thr Met Ile Asp Val
35 40 45

Gln Thr Thr Thr Val Gln Pro Ser Asp Glu Phe Glu Cys Pro Thr Arg
50 55 60

Phe Gly Tyr Phe Ala Asp Pro Lys Asp Pro Cys Lys Phe Tyr Ile Cys
65 70 75 80

Ser Asn Trp Glu Ala Ile His Lys Ser Cys Pro Gly Asn Thr Arg Trp
85 90 95

Asn Glu Lys Glu Leu Thr Cys Thr
100

<210> SEQ ID NO 62
<211> LENGTH: 131
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 13_isol

<400> SEQUENCE: 62

Met Ala Ser Ile Glu Gly Lys Tyr Lys Leu Glu Lys Ser Glu Lys Phe
1 5 10 15

Asp Glu Phe Leu Asp Lys Leu Gly Val Gly Phe Met Val Lys Thr Ala
20 25 30

Ala Lys Thr Leu Lys Pro Thr Phe Glu Val Ala Ile Glu Asn Asp Gln
35 40 45

Tyr Ile Phe Arg Ser Leu Ser Thr Phe Lys Asn Thr Glu Ala Lys Phe
50 55 60

Lys Leu Gly Glu Glu Phe Glu Glu Asp Arg Ala Asp Gly Lys Arg Val
65 70 75 80

Lys Thr Val Ile Gln Lys Glu Gly Asp Asn Lys Phe Val Gln Thr Gln
85 90 95

Phe Gly Asp Lys Glu Val Lys Ile Ile Arg Glu Phe Asn Gly Asp Glu
100 105 110

Val Val Val Thr Ala Ser Cys Asp Gly Val Thr Ser Val Arg Thr Tyr
115 120 125

Lys Arg Ile
130

<210> SEQ ID NO 63
<211> LENGTH: 213

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<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 7_isol

<400> SEQUENCE: 63

Met Met Lys Phe Leu Leu Ile Ala Ala Val Ala Phe Val Ala Val Ser
 1           5           10           15
Ala Asp Pro Ile His Tyr Asp Lys Ile Thr Glu Glu Ile Asn Lys Ala
 20           25           30
Ile Asp Asp Ala Ile Ala Ala Ile Glu Lys Ser Glu Thr Ile Asp Pro
 35           40           45
Met Lys Val Pro Asp His Thr Asp Lys Phe Glu Arg His Val Gly Ile
 50           55           60
Leu Asp Phe Lys Gly Glu Leu Ala Met Arg Asn Ile Glu Ala Arg Gly
 65           70           75           80
Leu Lys Gln Met Lys Arg Gln Gly Asp Ala Asn Val Lys Gly Glu Glu
 85           90           95
Gly Ile Val Lys Ala His Leu Leu Ile Gly Val His Asp Asp Ile Val
 100          105          110
Ser Met Glu Tyr Asp Leu Ala Tyr Lys Leu Gly Asp Leu His Pro Asn
 115          120          125
Thr His Val Ile Ser Asp Ile Gln Asp Phe Val Val Ala Leu Ser Leu
 130          135          140
Glu Ile Ser Asp Glu Gly Asn Ile Thr Met Thr Ser Phe Glu Val Arg
 145          150          155          160
Gln Phe Ala Asn Val Val Asn His Ile Gly Gly Leu Ser Ile Leu Asp
 165          170          175
Pro Ile Phe Gly Val Leu Ser Asp Val Leu Thr Ala Ile Phe Gln Asp
 180          185          190
Thr Val Arg Lys Glu Met Thr Lys Val Leu Ala Pro Ala Phe Lys Arg
 195          200          205

Glu Leu Glu Lys Asn
 210

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<210> SEQ ID NO 64
<211> LENGTH: 427
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 27_isol

<400> SEQUENCE: 64

Met Lys Phe Phe Leu Leu Ser Phe Val Leu Met Ile Val Ala Ala Thr
 1           5           10           15
Ala Thr Tyr Ala Ala His Val Gly Ser Gly Ser Arg Asp Asn Asn Asn
 20           25           30
Asn Lys Pro Val Pro Ala Glu Gly Phe Ala Lys Ala Ser Asn Glu Phe
 35           40           45
Gly Phe His Leu Leu Lys Glu Val Ile Gln His Arg Ser Ser Ser Gly
 50           55           60
Ser Arg Gly Ser Ser Glu Asn Val Leu Phe Ser Pro Tyr Ser Val Ala
 65           70           75           80
Val Ala Leu Ser Met Val His Gln Gly Thr Gln Gly Ser Thr Ala Glu
 85           90           95
Gln Phe Lys Arg Val Leu Tyr Tyr Asp Arg Val Gln Gln Leu Asn Gly
 100          105          110

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Met Lys Val Pro Asp His Thr Asp Lys Phe Glu Arg His Val Gly Ile
 50 55 60

Leu Asp Phe Lys Gly Glu Leu Ala Met Arg Asn Ile Glu Ala Arg Gly
 65 70 75 80

Leu Lys Gln Met Lys Arg Gln Gly Asp Ala Asn Val Lys Gly Glu Glu
 85 90 95

Gly Ile Val Lys Ala His Leu Leu Ile Gly Val His Asp Asp Ile Val
 100 105 110

Ser Met Glu Tyr Asp Leu Ala Tyr Lys Leu Gly Asp Leu His Pro Asn
 115 120 125

Thr His Val Ile Ser Asp Ile Gln Asp Phe Val Val Ala Leu Ser Leu
 130 135 140

Glu Ile Ser Asp Glu Gly Asn Ile Thr Met Thr Ser Phe Glu Val Arg
 145 150 155 160

Gln Phe Ala Asn Val Val Asn His Ile Gly Gly Leu Ser Ile Leu Asp
 165 170 175

Pro Ile Phe Gly Val Leu Ser Asp Val Leu Thr Ala Ile Phe Gln Asp
 180 185 190

Thr Val Arg Lys Glu Met Thr Lys Val Leu Ala Pro Ala Phe Lys Arg
 195 200 205

Glu Leu Glu Lys Asn
 210

<210> SEQ ID NO 66
 <211> LENGTH: 164
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 29_isol

<400> SEQUENCE: 66

Met Ala Leu Pro Arg Val Phe Phe Asp Ile Ala Ala Asp Asn Gln Pro
 1 5 10 15

Leu Gly Arg Ile Val Ile Glu Leu Arg Ser Asp Val Val Pro Lys Thr
 20 25 30

Ala Glu Asn Phe Arg Ala Leu Cys Thr Gly Glu Lys Gly Phe Gly Phe
 35 40 45

Lys Ser Ser Ser Phe His Arg Ile Ile Pro Asn Phe Met Ile Gln Gly
 50 55 60

Gly Asp Phe Thr Asn His Asn Gly Thr Gly Gly Lys Ser Ile Tyr Gly
 65 70 75 80

Asn Lys Phe Ala Asp Glu Asn Phe Thr Leu Gln His Thr Gly Pro Gly
 85 90 95

Ile Met Ser Met Ala Asn Ala Gly Pro Asn Thr Asn Gly Ser Gln Phe
 100 105 110

Phe Ile Thr Thr Val Lys Thr Thr Trp Leu Asp Gly Lys His Val Val
 115 120 125

Phe Gly Ser Val Val Glu Gly Met Asp Ile Val Lys Lys Val Glu Ser
 130 135 140

Tyr Gly Ser Gln Ser Gly Lys Pro Ser Lys Lys Val Thr Ile Ala Asn
 145 150 155 160

Cys Gly Gln Leu

<210> SEQ ID NO 67
 <211> LENGTH: 462

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<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 18_isol

<400> SEQUENCE: 67

Met Thr Arg Phe Ser Leu Thr Val Leu Ala Val Leu Ala Ala Cys Phe
 1          5          10          15
Gly Ser Asn Ile Arg Pro Asn Val Ala Thr Leu Glu Pro Lys Thr Val
 20          25          30
Cys Tyr Tyr Glu Ser Trp Val His Trp Arg Gln Gly Glu Gly Lys Met
 35          40          45
Asp Pro Glu Asp Ile Asp Thr Ser Leu Cys Thr His Ile Val Tyr Ser
 50          55          60
Tyr Phe Gly Ile Asp Ala Ala Thr His Glu Ile Lys Leu Leu Asp Glu
 65          70          75          80
Tyr Leu Met Lys Asp Leu His Asp Met Glu His Phe Thr Gln His Lys
 85          90          95
Gly Asn Ala Lys Ala Met Ile Ala Val Gly Gly Ser Thr Met Ser Asp
 100         105         110
Gln Phe Ser Lys Thr Ala Ala Val Glu His Tyr Arg Glu Thr Phe Val
 115         120         125
Val Ser Thr Val Asp Leu Met Thr Arg Tyr Gly Phe Asp Gly Val Met
 130         135         140
Ile Asp Trp Ser Gly Met Gln Ala Lys Asp Ser Asp Asn Phe Ile Lys
 145         150         155         160
Leu Leu Asp Lys Phe Asp Glu Lys Phe Ala His Thr Ser Phe Val Met
 165         170         175
Gly Val Thr Leu Pro Ala Thr Ile Ala Ser Tyr Asp Asn Tyr Asn Ile
 180         185         190
Pro Ala Ile Ser Asn Tyr Val Asp Phe Met Asn Val Leu Ser Leu Asp
 195         200         205
Tyr Thr Gly Ser Trp Ala His Thr Val Gly His Ala Ser Pro Phe Pro
 210         215         220
Glu Gln Leu Lys Thr Leu Glu Ala Tyr His Lys Arg Gly Ala Pro Arg
 225         230         235         240
His Lys Met Val Met Ala Val Pro Phe Tyr Ala Arg Thr Trp Ile Leu
 245         250         255
Glu Lys Met Asn Lys Gln Asp Ile Gly Asp Lys Ala Ser Gly Pro Gly
 260         265         270
Pro Arg Gly Gln Phe Thr Gln Thr Asp Gly Phe Leu Ser Tyr Asn Glu
 275         280         285
Leu Cys Val Gln Ile Gln Ala Glu Thr Asn Ala Phe Thr Ile Thr Arg
 290         295         300
Asp His Asp Asn Thr Ala Ile Tyr Ala Val Tyr Val His Ser Asn His
 305         310         315         320
Ala Glu Trp Ile Ser Phe Glu Asp Arg His Thr Leu Gly Glu Lys Ala
 325         330         335
Lys Asn Ile Thr Gln Gln Gly Tyr Ala Gly Met Ser Val Tyr Thr Leu
 340         345         350
Ser Asn Glu Asp Val His Gly Val Cys Gly Asp Lys Asn Pro Leu Leu
 355         360         365
His Ala Ile Gln Ser Asn Tyr Tyr His Gly Val Val Thr Glu Pro Thr
 370         375         380

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Ile Asp Ile Ser Phe Val Lys Asn Gly Pro Leu Ser Arg Ala Asp Leu
 290                295                300
Asp Thr Asn Asp Thr Phe Ile Val Glu Asn Gly Ser Glu Gly Leu Trp
 305                310                315
Val Trp Val Gly Lys Lys Ala Thr Gln Lys Glu Arg Gln Ser Ala Ile
                325                330                335
Lys Tyr Ala Met Glu Leu Ile Asn Lys Lys Lys Tyr Pro Asn Asn Thr
                340                345                350
Pro Val Thr Lys Val Leu Glu Gly Asp Glu Ser Val Glu Phe Lys Ser
                355                360                365
Leu Phe Glu Ser Trp Gln Met Ser Glu Gln Glu Lys Ile Thr Ser Ala
 370                375                380
Arg Leu Phe Arg Val Ser Arg Asn Gly Ile Phe Lys Gln Val Ala Asn
 385                390                395                400
Tyr Glu Pro Asp Asp Leu Glu Glu Asp Asn Ile Met Ile Leu Asp Val
                405                410                415
Met Asp Lys Ile Tyr Val Trp Ile Gly Asn Gln Phe Ala Glu Arg Ile
                420                425                430
Ala Asp Glu Ala His Val Asp Lys Val Ala Gln Arg Phe Ile Gln Glu
                435                440                445
Asp Lys Ser Gly Arg Lys Phe Arg Pro Asn Gln Ile Ile Lys Leu Lys
 450                455                460
Gln Gly Ser Glu Asp Gly Ala Phe Lys Ser Tyr Phe Pro Lys Trp Asn
 465                470                475                480

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<210> SEQ ID NO 69
<211> LENGTH: 213
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Der f 7_iso3

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<400> SEQUENCE: 69

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Met Met Lys Phe Leu Leu Ile Ala Ala Val Ala Phe Val Ala Val Ser
 1                5                10                15
Ala Asp Pro Ile His Tyr Asp Lys Ile Thr Glu Glu Ile Asn Lys Ala
                20                25                30
Ile Asp Asp Ala Ile Ala Ala Ile Glu Lys Ser Glu Thr Ile Asp Pro
                35                40                45
Met Lys Val Pro Asp His Thr Asp Lys Phe Glu Arg His Val Gly Ile
 50                55                60
Leu Asp Phe Lys Gly Glu Leu Ala Met Arg Asn Ile Glu Ala Arg Gly
 65                70                75                80
Leu Lys Gln Met Lys Arg Gln Gly Asp Ala Asn Val Lys Gly Glu Glu
                85                90                95
Gly Ile Val Lys Ala His Leu Leu Ile Gly Val His Asp Asp Ile Val
                100                105                110
Ser Met Glu Tyr Asp Leu Ala Tyr Lys Leu Gly Asp Leu His Pro Asn
                115                120                125
Thr His Val Ile Ser Asp Ile Gln Asp Phe Val Val Ala Leu Ser Leu
 130                135                140
Glu Ile Ser Asp Glu Gly Asn Ile Thr Met Thr Ser Phe Glu Val Arg
 145                150                155                160
Gln Phe Ala Asn Val Val Asn His Ile Gly Gly Leu Ser Ile Leu Asp
                165                170                175

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Glu Asp Arg Ser Lys Leu Lys Leu Gly Asp Pro Ala Lys Gly Met Ser
 290 295 300
 Pro Pro Gly Phe Ile Ser Gly Glu Glu Gly Val Leu Ser Tyr Ile Glu
 305 310 315 320
 Leu Cys Gln Leu Phe Gln Lys Glu Glu Trp His Ile Gln Tyr Asp Glu
 325 330 335
 Tyr Tyr Asn Ala Pro Tyr Gly Tyr Asn Asp Lys Ile Trp Val Gly Tyr
 340 345 350
 Asp Asp Leu Ala Ser Ile Ser Cys Lys Leu Ala Phe Leu Lys Glu Leu
 355 360 365
 Gly Val Ser Gly Val Ile Val Trp Ser Leu Glu Asn Asp Asp Phe Lys
 370 375 380
 Gly His Cys Gly Pro Lys Asn Pro Leu Leu Asn Lys Val His Asn Met
 385 390 395 400
 Ile Asn Gly Asp Glu Lys Asn Ser Phe Glu Cys Ile Leu Gly Pro Ser
 405 410 415
 Thr Thr Thr Pro Thr Pro Thr Thr Thr Pro Thr Thr Thr Pro Thr Pro
 420 425 430
 Ser Pro Thr Thr Pro Thr Pro Ser Pro Thr Thr Pro Thr Thr Thr Pro
 435 440 445
 Ser Pro Thr Thr Pro Thr Pro Ser Pro Thr Thr Pro Thr Thr Thr Pro
 450 455 460
 Ser Pro Thr Thr Pro Thr Pro Thr Thr Pro Thr Pro Thr Pro Thr Thr
 465 470 475 480
 Ser Thr Pro Ser Pro Thr Thr Thr Glu His Thr Ser Glu Thr Pro Lys
 485 490 495
 Tyr Thr Thr Tyr Val Asp Gly His Leu Ile Lys Cys Tyr Lys Glu Gly
 500 505 510
 Asp Ile Pro His Pro Thr Asn Ile His Lys Tyr Leu Val Cys Glu Phe
 515 520 525
 Val Asn Gly Gly Trp Trp Val His Ile Met Pro Cys Pro Pro Gly Thr
 530 535 540
 Ile Trp Cys Gln Glu Lys Leu Thr Cys Ile Gly Glu
 545 550 555

<210> SEQ ID NO 73
 <211> LENGTH: 356
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 20_iso2

<400> SEQUENCE: 73

Met Val Asp Gln Ala Thr Leu Ser Lys Leu Glu Ala Gly Phe Gln Lys
 1 5 10 15
 Leu Gln Asn Ala Gln Asp Cys His Ser Leu Leu Lys Lys Tyr Leu Thr
 20 25 30
 Arg Asp Val Leu Asp Gln Leu Lys Thr Lys Lys Thr Asp Met Gly Ala
 35 40 45
 Thr Leu Leu Asp Val Ile Gln Ser Gly Val Glu Asn Leu Asp Ser Gly
 50 55 60
 Val Gly Ile Tyr Ala Pro Asp Ala Gln Ser Tyr Lys Thr Phe Ala Ala
 65 70 75 80
 Leu Phe Asp Pro Ile Ile Asp Asp Tyr His Lys Gly Phe Lys Pro Thr
 85 90 95

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Asp Lys His Pro Gln Thr Asp Phe Gly Asn Ile Glu His Phe Val Asn
 100 105 110
 Val Asp Pro Lys Asn Glu Tyr Val Ile Ser Thr Arg Val Arg Cys Gly
 115 120 125
 Arg Ser Leu Lys Gly Tyr Pro Phe Asn Pro Met Leu Thr Glu Ala Gln
 130 135 140
 Tyr Lys Glu Met Glu Thr Lys Val Lys Gly Gln Leu Ala Thr Phe Glu
 145 150 155 160
 Gly Glu Leu Lys Gly Thr Tyr Tyr Pro Leu Leu Gly Met Asp Lys Ala
 165 170 175
 Thr Gln Gln Lys Leu Ile Asp Asp His Phe Leu Phe Lys Glu Gly Asp
 180 185 190
 Arg Phe Leu Gln Ala Ala Asn Ala Cys Arg Tyr Trp Pro Val Gly Arg
 195 200 205
 Gly Ile Phe His Asn Asp Lys Lys Thr Phe Leu Met Trp Val Asn Glu
 210 215 220
 Glu Asp His Leu Arg Ile Ile Ser Met Gln Lys Gly Gly Asp Leu Lys
 225 230 235 240
 Glu Val Phe Gly Arg Leu Val Lys Ala Val Lys His Ile Glu Gln Lys
 245 250
 Ile Pro Phe Ser Arg Asp Asp Arg Leu Gly Tyr Leu Thr Phe Cys Pro
 260 265 270
 Thr Asn Leu Gly Thr Thr Ile Arg Ala Ser Val His Ile Lys Leu Pro
 275 280 285
 Lys Leu Ala Ala Asp Arg Lys Lys Leu Glu Glu Val Ala Ala Arg Tyr
 290 295 300
 Asn Leu Gln Val Arg Gly Thr Ala Gly Glu His Thr Glu Ser Val Gly
 305 310 315 320
 Gly Ile Tyr Asp Ile Ser Asn Lys Arg Arg Met Gly Leu Thr Glu Tyr
 325 330 335
 Gln Ala Val Lys Glu Met Gln Asp Gly Ile Ile Glu Leu Ile Lys Met
 340 345 350
 Glu Lys Ser Leu
 355

<210> SEQ ID NO 74

<211> LENGTH: 77

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Der f 1_iso1_TI

<400> SEQUENCE: 74

Phe Trp Pro Thr Val Thr Arg Leu Trp Ile Phe Leu Asn Arg Asn Ser
 1 5 10 15
 Ser Ile Ala His Leu Asn Thr Asp Val Thr Ala Ile Gln Tyr Gln Glu
 20 25 30
 Ala Ser Asn Thr Ser Asn Lys Met Val Ser Leu Lys Lys Glu Ala Ile
 35 40 45
 His Thr Leu His Glu Asn Asn Asn Ala Asp Asp Gln Ile Arg Asn Ile
 50 55 60
 Thr Val Ser Gln Thr Thr Ala Lys Phe Ile His Gln Met
 65 70 75

<210> SEQ ID NO 75

<211> LENGTH: 61

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<212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 1_iso1_TI

<400> SEQUENCE: 75

Ser Asn Ile Met Met Asp Glu Gln Ser Phe Asn Met Thr Met Val Ile
 1 5 10 15

Asn Gln Thr Ile Met Pro Ser Thr Leu Ser Val Thr Glu Val His Lys
 20 25 30

Ala Ser Ile Ile Gly Ser Tyr Glu Thr Val Gly Ile Gln Pro Gly Val
 35 40 45

Ile Ala Asp Thr Asp Ile Ser Lys Pro Glu Thr Thr Ser
 50 55 60

<210> SEQ ID NO 76
 <211> LENGTH: 48
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 1_iso1_TI

<400> SEQUENCE: 76

Met Val Ile Asn Gln Thr Ile Met Pro Ser Thr Leu Ser Val Thr Glu
 1 5 10 15

Val His Lys Ala Ser Ile Ile Gly Ser Tyr Glu Thr Val Gly Ile Gln
 20 25 30

Pro Gly Val Ile Ala Asp Thr Asp Ile Ser Lys Pro Glu Thr Thr Ser
 35 40 45

<210> SEQ ID NO 77
 <211> LENGTH: 26
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 1_iso1_TI

<400> SEQUENCE: 77

Ile Gly Ser Tyr Glu Thr Val Gly Ile Gln Pro Gly Val Ile Ala Asp
 1 5 10 15

Thr Asp Ile Ser Lys Pro Glu Thr Thr Ser
 20 25

<210> SEQ ID NO 78
 <211> LENGTH: 77
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 1_iso2_TI

<400> SEQUENCE: 78

Phe Trp Pro Thr Val Thr Arg Leu Trp Ile Phe Leu Asn Arg Asn Ser
 1 5 10 15

Ser Ile Ala His Leu Asn Thr Asp Val Thr Ala Ile Gln Tyr Gln Glu
 20 25 30

Ala Ser Asn Thr Ser Asn Lys Met Val Ser Leu Lys Lys Glu Ala Ile
 35 40 45

His Thr Leu His Glu Asn Asn Asn Ala Asp Asp Gln Ile Arg Asn Ile
 50 55 60

Thr Val Ser Gln Thr Thr Ala Lys Phe Ile His Gln Met
 65 70 75

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<210> SEQ ID NO 79
 <211> LENGTH: 61
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 1_iso2_TI

 <400> SEQUENCE: 79

 Ser Asn Ile Met Met Asp Glu Gln Ser Phe Asn Met Thr Met Val Ile
 1 5 10 15

 Asn Gln Thr Ile Met Pro Ser Thr Leu Ser Val Thr Glu Val His Lys
 20 25 30

 Ala Ser Ile Ile Gly Ser Tyr Glu Thr Val Gly Ile Gln Pro Gly Val
 35 40 45

 Ile Ala Asp Thr Asp Ile Ser Lys Pro Glu Thr Thr Ser
 50 55 60

<210> SEQ ID NO 80
 <211> LENGTH: 48
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 1_iso2_TI

 <400> SEQUENCE: 80

 Met Val Ile Asn Gln Thr Ile Met Pro Ser Thr Leu Ser Val Thr Glu
 1 5 10 15

 Val His Lys Ala Ser Ile Ile Gly Ser Tyr Glu Thr Val Gly Ile Gln
 20 25 30

 Pro Gly Val Ile Ala Asp Thr Asp Ile Ser Lys Pro Glu Thr Thr Ser
 35 40 45

<210> SEQ ID NO 81
 <211> LENGTH: 26
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 1_iso2_TI

 <400> SEQUENCE: 81

 Ile Gly Ser Tyr Glu Thr Val Gly Ile Gln Pro Gly Val Ile Ala Asp
 1 5 10 15

 Thr Asp Ile Ser Lys Pro Glu Thr Thr Ser
 20 25

<210> SEQ ID NO 82
 <211> LENGTH: 23
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 23_iso1_TI

 <400> SEQUENCE: 82

 Ile Phe Val Gln Ile Gly Lys Leu Tyr Ile Lys Val Val Gln Val Ile
 1 5 10 15

 Gln Asp Gly Met Lys Lys Asn
 20

<210> SEQ ID NO 83
 <211> LENGTH: 25
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:

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<223> OTHER INFORMATION: Der f 11_isol_TI

<400> SEQUENCE: 83

Tyr Val Ile Asn Val Ile Asn Trp His Val Lys Thr Arg Asn Leu Gln
1 5 10 15

Thr Ile Leu Pro Lys Leu Asn His Asn
20 25

<210> SEQ ID NO 84

<211> LENGTH: 28

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Der f 11_isol_TI

<400> SEQUENCE: 84

Thr Met Lys Tyr Ile Lys Asn Leu Glu Phe Leu Met Asn Glu Tyr Arg
1 5 10 15

Asn Leu Gln Leu Asn Ser Asn Leu Lys Ile Cys
20 25

<210> SEQ ID NO 85

<211> LENGTH: 37

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Der f 15_isol_TI

<400> SEQUENCE: 85

Trp Ile Gly Ser Ile Leu Asp Leu Asp Trp Val Thr Arg Lys Ser Thr
1 5 10 15

Asn Lys Thr Ile Trp Leu Trp Leu Glu Asn Leu Lys Thr Leu Leu Asn
20 25 30

Leu Met Ala Thr Cys
35

<210> SEQ ID NO 86

<211> LENGTH: 15

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Der f 32_isol_TI

<400> SEQUENCE: 86

Leu Leu Lys Lys Ser Leu Lys Lys His Ile Asn Ile Gly Lys Lys
1 5 10 15

<210> SEQ ID NO 87

<211> LENGTH: 44

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Der f 25_isol_TI

<400> SEQUENCE: 87

Gly Pro Leu Val Leu Val Lys Gln Pro Val His Asn Lys His Lys Lys
1 5 10 15

Phe Ile Lys Asn Phe Asp Asn Gly Phe Leu Lys Met Phe His His Lys
20 25 30

Leu Pro Lys Gln Phe Glu Ser Phe Met Val Val Gln
35 40

<210> SEQ ID NO 88

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<211> LENGTH: 17
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 25_iso1_TI

<400> SEQUENCE: 88

Met Phe His His Lys Leu Pro Lys Gln Phe Glu Ser Phe Met Val Val
 1 5 10 15

Gln

<210> SEQ ID NO 89
 <211> LENGTH: 15
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 16_iso1_TI

<400> SEQUENCE: 89

Tyr Lys Arg Ile Lys Val Ala Val Asn Phe Asp Gln Ile Arg Leu
 1 5 10 15

<210> SEQ ID NO 90
 <211> LENGTH: 27
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 26_iso1_TI

<400> SEQUENCE: 90

Glu Ile Ser Val His Phe Ala Leu Val Lys Lys Asp Leu Val Leu Asn
 1 5 10 15

His Pro His Phe Ile Val Ser Tyr Pro Ile Leu
 20 25

<210> SEQ ID NO 91
 <211> LENGTH: 16
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 26_iso1_TI

<400> SEQUENCE: 91

Lys Arg Trp Lys Ala Met Ala His Asn Arg Val Asn His Pro Arg Lys
 1 5 10 15

<210> SEQ ID NO 92
 <211> LENGTH: 23
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 23_iso2_TI

<400> SEQUENCE: 92

Ile Phe Val Gln Ile Gly Lys Leu Tyr Ile Lys Val Val Gln Val Ile
 1 5 10 15

Gln Asp Gly Met Lys Lys Asn
 20

<210> SEQ ID NO 93
 <211> LENGTH: 15
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 13_iso1_TI

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<400> SEQUENCE: 93

Lys Val Thr Ile Asn Leu Phe Lys His Asn Ser Val Ile Lys Lys
 1 5 10 15

<210> SEQ ID NO 94

<211> LENGTH: 17

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Der f 7_iso1_TI

<400> SEQUENCE: 94

Leu Pro Ile Asn Ser Asn Val Met Leu Val Phe Trp Ile Ser Lys Val
 1 5 10 15

Asn

<210> SEQ ID NO 95

<211> LENGTH: 46

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Der f 27_iso1_TI

<400> SEQUENCE: 95

Met Ile Leu Thr Lys Phe Val Asn Gln Ser Ile Gln Leu Leu Leu Met
 1 5 10 15

Lys Leu Leu Asp Lys Leu Leu Val Val Ser His Gln Leu Phe Ile Ser
 20 25 30

Arg Lys Leu Asn Ser Val His His Ile Asn Cys Arg Lys Phe
 35 40 45

<210> SEQ ID NO 96

<211> LENGTH: 17

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Der f 7_iso2_TI

<400> SEQUENCE: 96

Leu Pro Ile Asn Ser Asn Val Met Leu Val Phe Trp Ile Ser Lys Val
 1 5 10 15

Asn

<210> SEQ ID NO 97

<211> LENGTH: 16

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Der f 29_iso1_TI

<400> SEQUENCE: 97

Lys Arg Trp Lys Ala Met Ala His Asn Arg Val Asn His Pro Arg Lys
 1 5 10 15

<210> SEQ ID NO 98

<211> LENGTH: 15

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Der f 18_iso1_TI

<400> SEQUENCE: 98

Lys Pro Cys Ile Ile Val Pro Ile Thr Tyr Arg His Leu Thr Lys
 1 5 10 15

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<210> SEQ ID NO 99
 <211> LENGTH: 15
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 16_iso2_TI

<400> SEQUENCE: 99

Tyr Lys Arg Ile Lys Val Ala Val Asn Phe Asp Gln Ile Arg Leu
 1 5 10 15

<210> SEQ ID NO 100
 <211> LENGTH: 17
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 7_iso3_TI

<400> SEQUENCE: 100

Leu Pro Ile Asn Ser Asn Val Met Leu Val Phe Trp Ile Ser Lys Val
 1 5 10 15

Asn

<210> SEQ ID NO 101
 <211> LENGTH: 30
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 20_iso1_TI

<400> SEQUENCE: 101

Cys Val Met Trp Ile Gln Ile Met Asn Leu Ser Phe Gln His Val Tyr
 1 5 10 15

Val Val Ala Asp His Cys Lys Val Ile His Leu Ile His Ala
 20 25 30

<210> SEQ ID NO 102
 <211> LENGTH: 52
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 20_isol_TI

<400> SEQUENCE: 102

Trp Lys Lys Leu Leu Ala Asn Ile Ile Tyr Lys Tyr Val Val Leu Pro
 1 5 10 15

Val Asn Thr Pro Lys Val Leu Ala Val Phe Thr Ile Ser Val Ile Asn
 20 25 30

Val Val Trp Val Leu Leu Asn Ile Arg Pro Ser Lys Arg Cys Lys Met
 35 40 45

Val Phe Leu Asn
 50

<210> SEQ ID NO 103
 <211> LENGTH: 36
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 20_isol_TI

<400> SEQUENCE: 103

Val Asn Thr Pro Lys Val Leu Ala Val Phe Thr Ile Ser Val Ile Asn
 1 5 10 15

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Val Val Trp Val Leu Leu Asn Ile Arg Pro Ser Lys Arg Cys Lys Met
 20 25 30

Val Phe Leu Asn
 35

<210> SEQ ID NO 104
 <211> LENGTH: 26
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 20_iso1_TI

<400> SEQUENCE: 104

Thr Ile Ser Val Ile Asn Val Val Trp Val Leu Leu Asn Ile Arg Pro
 1 5 10 15

Ser Lys Arg Cys Lys Met Val Phe Leu Asn
 20 25

<210> SEQ ID NO 105
 <211> LENGTH: 18
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 1_iso3_TI

<400> SEQUENCE: 105

Glu Gln Val Met Val Leu Leu Asn Thr Glu Pro Asp Met Val Ser Leu
 1 5 10 15

Ala Val

<210> SEQ ID NO 106
 <211> LENGTH: 42
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 1_iso3_TI

<400> SEQUENCE: 106

Val Ala His Met Asp His His Ile Gln Tyr Asn Gln Pro Ala Thr Glu
 1 5 10 15

Leu Val Met Val Thr Leu Pro Ile Ala Val Met Pro Thr Gln Ser Glu
 20 25 30

Lys Asn Lys Leu Val Leu Pro Phe Ser Phe
 35 40

<210> SEQ ID NO 107
 <211> LENGTH: 37
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 15_iso2_TI

<400> SEQUENCE: 107

Trp Ile Gly Ser Ile Leu Asp Leu Asp Trp Val Thr Arg Lys Ser Thr
 1 5 10 15

Asn Lys Thr Ile Trp Leu Trp Leu Glu Asn Leu Lys Thr Leu Leu Asn
 20 25 30

Leu Met Ala Thr Cys
 35

<210> SEQ ID NO 108
 <211> LENGTH: 29

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<212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 20_iso2_TI

 <400> SEQUENCE: 108

 Val Val Ser Met Ile Leu Val Thr Asn Asp Glu Trp Val Ser Pro Asn
 1 5 10 15

 Thr Lys Leu Leu Arg Lys Cys Lys Met Ala Ser Leu Asn
 20 25

 <210> SEQ ID NO 109
 <211> LENGTH: 22
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Der f 20_iso2_TI

 <400> SEQUENCE: 109

 Thr Asn Asp Glu Trp Val Ser Pro Asn Thr Lys Leu Leu Arg Lys Cys
 1 5 10 15

 Lys Met Ala Ser Leu Asn
 20

 <210> SEQ ID NO 110
 <211> LENGTH: 56
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref1-g11-id30

 <400> SEQUENCE: 110

 Arg Lys Ser Asn Val Thr Arg Ile His Met Asp Gly Thr Arg Thr Ala
 1 5 10 15

 Leu Val Arg Ile Arg Thr Ala Leu Val Arg Thr Arg Thr Asp Val Ile
 20 25 30

 Arg Thr Ala Leu Val His Met Ile Gly Glu Ala Leu Asp Leu Leu Ser
 35 40 45

 Thr Lys Arg Gly Val Ala Met Ser
 50 55

 <210> SEQ ID NO 111
 <211> LENGTH: 40
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref1-g11-id31

 <400> SEQUENCE: 111

 Leu Val Arg Ile Arg Thr Ala Leu Val Arg Thr Arg Thr Asp Val Ile
 1 5 10 15

 Arg Thr Ala Leu Val His Met Ile Gly Glu Ala Leu Asp Leu Leu Ser
 20 25 30

 Thr Lys Arg Gly Val Ala Met Ser
 35 40

 <210> SEQ ID NO 112
 <211> LENGTH: 38
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref1-g11-id32

 <400> SEQUENCE: 112

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Arg Ile Arg Thr Ala Leu Val Arg Thr Arg Thr Asp Val Ile Arg Thr
 1 5 10 15

Ala Leu Val His Met Ile Gly Glu Ala Leu Asp Leu Leu Ser Thr Lys
 20 25 30

Arg Gly Val Ala Met Ser
 35

<210> SEQ ID NO 113
 <211> LENGTH: 21
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref1-g11-id33

<400> SEQUENCE: 113

Leu Val His Met Ile Gly Glu Ala Leu Asp Leu Leu Ser Thr Lys Arg
 1 5 10 15

Gly Val Ala Met Ser
 20

<210> SEQ ID NO 114
 <211> LENGTH: 10
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref1-g11-id34

<400> SEQUENCE: 114

Leu Ser Thr Lys Arg Gly Val Ala Met Ser
 1 5 10

<210> SEQ ID NO 115
 <211> LENGTH: 10
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref1-g12-id35

<400> SEQUENCE: 115

Lys Gly Ala Cys Ala Arg His Cys Asn Arg
 1 5 10

<210> SEQ ID NO 116
 <211> LENGTH: 9
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref1-g12-id36

<400> SEQUENCE: 116

Gly Ala Cys Ala Arg His Cys Asn Arg
 1 5

<210> SEQ ID NO 117
 <211> LENGTH: 76
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref2-g13-id37

<400> SEQUENCE: 117

Lys Gly Ala Cys Ala Arg His Cys Asn Arg Ser Trp Arg Thr Arg Ala
 1 5 10 15

Ile Gly Cys Arg Gly Gly Asn Arg Ser Asn Ser Ser Arg Gly Ser Ser

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20	25	30
Gly Thr Cys Leu Asn Ser Ala Ala Leu Gly His His Ser Val Ala Thr		
35	40	45
Trp Thr Ser Lys Val Ala Ala Glu Thr Asp Thr Lys His Leu Ser Gln		
50	55	60
Lys Lys Lys Arg Lys Glu Lys Lys Ile Ala Tyr Ile		
65	70	75

<210> SEQ ID NO 118
 <211> LENGTH: 75
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref2-g13-id38

<400> SEQUENCE: 118

Gly Ala Cys Ala Arg His Cys Asn Arg Ser Trp Arg Thr Arg Ala Ile		
1	5	10
Gly Cys Arg Gly Gly Asn Arg Ser Asn Ser Ser Arg Gly Ser Ser Gly		
20	25	30
Thr Cys Leu Asn Ser Ala Ala Leu Gly His His Ser Val Ala Thr Trp		
35	40	45
Thr Ser Lys Val Ala Ala Glu Thr Asp Thr Lys His Leu Ser Gln Lys		
50	55	60
Lys Lys Arg Lys Glu Lys Lys Ile Ala Tyr Ile		
65	70	75

<210> SEQ ID NO 119
 <211> LENGTH: 49
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref2-g13-id39

<400> SEQUENCE: 119

Ser Arg Gly Ser Ser Gly Thr Cys Leu Asn Ser Ala Ala Leu Gly His		
1	5	10
His Ser Val Ala Thr Trp Thr Ser Lys Val Ala Ala Glu Thr Asp Thr		
20	25	30
Lys His Leu Ser Gln Lys Lys Lys Arg Lys Glu Lys Lys Ile Ala Tyr		
35	40	45

Ile

<210> SEQ ID NO 120
 <211> LENGTH: 33
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref2-g13-id40

<400> SEQUENCE: 120

His Ser Val Ala Thr Trp Thr Ser Lys Val Ala Ala Glu Thr Asp Thr		
1	5	10
Lys His Leu Ser Gln Lys Lys Lys Arg Lys Glu Lys Lys Ile Ala Tyr		
20	25	30

Ile

<210> SEQ ID NO 121
 <211> LENGTH: 87
 <212> TYPE: PRT

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<210> SEQ ID NO 124
 <211> LENGTH: 87
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref1-g11-id33

 <400> SEQUENCE: 124

 Met Ala Arg Gln Gln Trp Glu Leu Gln Gly Asp Arg Arg Cys Gln Ser
 1 5 10 15

 Gln Leu Glu Arg Ala Asn Leu Arg Pro Cys Glu Gln His Leu Met Gln
 20 25 30

 Lys Ile Gln Arg Asp Glu Asp Ser Tyr Gly Arg Asp Pro Tyr Ser Pro
 35 40 45

 Ser Gln Asp Pro Tyr Ser Pro Ser Gln Asp Pro Asp Arg Arg Asp Pro
 50 55 60

 Tyr Ser Leu Val His Met Ile Gly Glu Ala Leu Asp Leu Leu Ser Thr
 65 70 75 80

 Lys Arg Gly Val Ala Met Ser
 85

<210> SEQ ID NO 125
 <211> LENGTH: 87
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref1-g11-id34

 <400> SEQUENCE: 125

 Met Ala Arg Gln Gln Trp Glu Leu Gln Gly Asp Arg Arg Cys Gln Ser
 1 5 10 15

 Gln Leu Glu Arg Ala Asn Leu Arg Pro Cys Glu Gln His Leu Met Gln
 20 25 30

 Lys Ile Gln Arg Asp Glu Asp Ser Tyr Gly Arg Asp Pro Tyr Ser Pro
 35 40 45

 Ser Gln Asp Pro Tyr Ser Pro Ser Gln Asp Pro Asp Arg Arg Asp Pro
 50 55 60

 Tyr Ser Pro Ser Pro Tyr Asp Arg Arg Gly Ala Gly Ser Leu Ser Thr
 65 70 75 80

 Lys Arg Gly Val Ala Met Ser
 85

<210> SEQ ID NO 126
 <211> LENGTH: 104
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref1-g12-id35

 <400> SEQUENCE: 126

 Met Ala Arg Gln Gln Trp Glu Leu Gln Gly Asp Arg Arg Cys Gln Ser
 1 5 10 15

 Gln Leu Glu Arg Ala Asn Leu Arg Pro Cys Glu Gln His Leu Met Gln
 20 25 30

 Lys Ile Gln Arg Asp Glu Asp Ser Tyr Gly Arg Asp Pro Tyr Ser Pro
 35 40 45

 Ser Gln Asp Pro Tyr Ser Pro Ser Gln Asp Pro Asp Arg Arg Asp Pro
 50 55 60

-continued

Tyr Ser Pro Ser Pro Tyr Asp Arg Arg Gly Ala Gly Ser Ser Gln His
65 70 75 80

Gln Glu Arg Cys Cys Asn Glu Leu Asn Glu Phe Glu Asn Asn Lys Gly
85 90 95

Ala Cys Ala Arg His Cys Asn Arg
100

<210> SEQ ID NO 127
 <211> LENGTH: 104
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref1-gl2-id36

<400> SEQUENCE: 127

Met Ala Arg Gln Gln Trp Glu Leu Gln Gly Asp Arg Arg Cys Gln Ser
1 5 10 15

Gln Leu Glu Arg Ala Asn Leu Arg Pro Cys Glu Gln His Leu Met Gln
20 25 30

Lys Ile Gln Arg Asp Glu Asp Ser Tyr Gly Arg Asp Pro Tyr Ser Pro
35 40 45

Ser Gln Asp Pro Tyr Ser Pro Ser Gln Asp Pro Asp Arg Arg Asp Pro
50 55 60

Tyr Ser Pro Ser Pro Tyr Asp Arg Arg Gly Ala Gly Ser Ser Gln His
65 70 75 80

Gln Glu Arg Cys Cys Asn Glu Leu Asn Glu Phe Glu Asn Asn Gln Gly
85 90 95

Ala Cys Ala Arg His Cys Asn Arg
100

<210> SEQ ID NO 128
 <211> LENGTH: 158
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref2-gl3-id37

<400> SEQUENCE: 128

Met Ala Arg Gln Gln Trp Glu Leu Gln Gly Asp Arg Arg Cys Gln Ser
1 5 10 15

Gln Leu Glu Arg Ala Asn Leu Arg Pro Cys Glu Gln His Leu Met Gln
20 25 30

Lys Ile Gln Arg Asp Glu Asp Ser Tyr Glu Arg Asp Pro Tyr Ser Pro
35 40 45

Ser Gln Asp Pro Tyr Ser Pro Ser Pro Tyr Asp Arg Arg Gly Ala Gly
50 55 60

Ser Ser Gln His Gln Glu Arg Cys Cys Asn Glu Leu Asn Glu Phe Glu
65 70 75 80

Asn Asn Lys Gly Ala Cys Ala Arg His Cys Asn Arg Ser Trp Arg Thr
85 90 95

Arg Ala Ile Gly Cys Arg Gly Gly Asn Arg Ser Asn Ser Ser Arg Gly
100 105 110

Ser Ser Gly Thr Cys Leu Asn Ser Ala Ala Leu Gly His His Ser Val
115 120 125

Ala Thr Trp Thr Ser Lys Val Ala Ala Glu Thr Asp Thr Lys His Leu
130 135 140

Ser Gln Lys Lys Lys Arg Lys Glu Lys Lys Ile Ala Tyr Ile
145 150 155

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<210> SEQ ID NO 129
 <211> LENGTH: 158
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref2-g13-id38

<400> SEQUENCE: 129

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Met Ala Arg Gln Gln Trp Glu Leu Gln Gly Asp Arg Arg Cys Gln Ser
1           5           10           15
Gln Leu Glu Arg Ala Asn Leu Arg Pro Cys Glu Gln His Leu Met Gln
20           25           30
Lys Ile Gln Arg Asp Glu Asp Ser Tyr Glu Arg Asp Pro Tyr Ser Pro
35           40           45
Ser Gln Asp Pro Tyr Ser Pro Ser Pro Tyr Asp Arg Arg Gly Ala Gly
50           55           60
Ser Ser Gln His Gln Glu Arg Cys Cys Asn Glu Leu Asn Glu Phe Glu
65           70           75           80
Asn Asn Gln Gly Ala Cys Ala Arg His Cys Asn Arg Ser Trp Arg Thr
85           90           95
Arg Ala Ile Gly Cys Arg Gly Gly Asn Arg Ser Asn Ser Ser Arg Gly
100          105          110
Ser Ser Gly Thr Cys Leu Asn Ser Ala Ala Leu Gly His His Ser Val
115          120          125
Ala Thr Trp Thr Ser Lys Val Ala Ala Glu Thr Asp Thr Lys His Leu
130          135          140
Ser Gln Lys Lys Lys Arg Lys Glu Lys Lys Ile Ala Tyr Ile
145          150          155
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<210> SEQ ID NO 130
 <211> LENGTH: 158
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Arah2-ref2-g13-id39

<400> SEQUENCE: 130

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Met Ala Arg Gln Gln Trp Glu Leu Gln Gly Asp Arg Arg Cys Gln Ser
1           5           10           15
Gln Leu Glu Arg Ala Asn Leu Arg Pro Cys Glu Gln His Leu Met Gln
20           25           30
Lys Ile Gln Arg Asp Glu Asp Ser Tyr Glu Arg Asp Pro Tyr Ser Pro
35           40           45
Ser Gln Asp Pro Tyr Ser Pro Ser Pro Tyr Asp Arg Arg Gly Ala Gly
50           55           60
Ser Ser Gln His Gln Glu Arg Cys Cys Asn Glu Leu Asn Glu Phe Glu
65           70           75           80
Asn Asn Gln Arg Cys Met Cys Glu Ala Leu Gln Gln Ile Met Glu Asn
85           90           95
Gln Ser Asp Arg Leu Gln Gly Arg Gln Gln Glu Gln Gln Ser Arg Gly
100          105          110
Ser Ser Gly Thr Cys Leu Asn Ser Ala Ala Leu Gly His His Ser Val
115          120          125
Ala Thr Trp Thr Ser Lys Val Ala Ala Glu Thr Asp Thr Lys His Leu
130          135          140
Ser Gln Lys Lys Lys Arg Lys Glu Lys Lys Ile Ala Tyr Ile
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-continued

Val Glu Ser Gly Gly Arg Asp Arg Tyr
145 150

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<212> TYPE: PRT
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Gln Leu Glu Arg Ala Asn Leu Arg Pro Cys Glu Gln His Leu Met Gln
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Lys Ile Gln Arg Asp Glu Asp Ser Tyr Glu Arg Asp Pro Tyr Ser Pro
35 40 45
Ser Gln Asp Pro Tyr Ser Pro Ser Pro Tyr Asp Arg Arg Gly Ala Gly
50 55 60
Ser Ser Gln His Gln Glu Arg Cys Cys Asn Glu Leu Asn Glu Phe Glu
65 70 75 80
Asn Asn Gln Arg Cys Met Cys Glu Ala Leu Gln Gln Ile Met Glu Asn
85 90 95
Gln Ser Asp Arg Leu Gln Gly Arg Gln Gln Glu Gln Gln Phe Lys Arg
100 105 110
Glu Leu Arg Asn Leu Pro Gln Gln Cys Gly Leu Arg Ala Pro Gln Arg
115 120 125
Cys Asp Leu Asp Val Glu Ser Gly Gly Arg Asp Arg Tyr
130 135 140

The invention claimed is:

1. A method for reducing immunogenicity or allergenicity of a composition, the method comprising treating the composition to remove cationic proteins, wherein the removed cationic proteins comprise proteins resulting from transcription infidelity.

2. The method of claim 1, which comprises removing at least 50% by weight of the cationic proteins of said composition having an isoelectric point above 7.4.

3. The method of claim 1, wherein the treated composition contains less than 2% by weight of cationic proteins having an isoelectric point above 8.

4. The method of claim 1, wherein the removed cationic proteins comprise a peptide sequence resulting from a transcription infidelity deletion.

5. The method of claim 1, wherein the composition is treated by cation exchange and/or affinity chromatography.

6. The method of claim 5, which comprises (i) providing a solution of the composition, (ii) adjusting, if necessary, the solution of the composition to have a pH between 7 and 9, (iii) subjecting the solution to cation exchange chromatography allowing binding of components of the solution having an isoelectric point above the adjusted pH, and (iv) recovering the unbound flow-through.

7. The method of claim 6, comprising a further step of subjecting the composition to affinity chromatography using transcription infidelity antibodies.

8. The method of claim 1, wherein the composition is or comprises a food ingredient, a feed ingredient, or a drug.

9. The method of claim 1, wherein the composition comprises purified and/or recombinant proteins, polypeptides or peptides.

10. The method of claim 1, wherein the composition comprises milk or a dairy product and the method comprises removing at least one protein comprising a transcription infidelity peptide sequence selected from SEQ ID NO: 1 to 5.

11. The method of claim 1, wherein the composition comprises a peanut product, and the method comprises removing at least one protein comprising a transcription infidelity peptide sequence selected from SEQ ID NO: 110 to 120.

12. A method for preparing a food product comprising (i) providing a food product preparation containing cationic proteins, (ii) treating the food product preparation to remove cationic proteins comprising proteins resulting from transcription infidelity, and (iii) optionally formulating the treated food product with one or more suitable excipients.

13. The method of claim 12, wherein the treated food product contains less than 2% by weight of cationic proteins resulting from transcription infidelity.

14. A method for preparing a pharmaceutical product comprising (i) providing a pharmaceutical product preparation, (ii) treating the pharmaceutical product preparation to remove cationic proteins comprising proteins resulting from transcription infidelity, and (iii) optionally formulating the treated pharmaceutical product with one or more suitable excipients.

15. The method of claim 14, wherein the pharmaceutical product is or comprises a vaccine, an immunogen, an allergen or a drug.

16. A method for reducing immunogenicity or allergenicity of a composition, the method comprising treating the

composition to remove cationic proteins, and wherein the method comprises (i) providing a solution of the composition, (ii) adjusting, if necessary, the solution of the composition to have a pH between 7 and 9, (iii) subjecting the solution to cation exchange chromatography allowing binding of components of the solution having an isoelectric point above the adjusted pH, (iv) recovering the unbound flow-through, and (v) subjecting the flow-through to affinity chromatography using transcription infidelity antibodies. 5

17. A method for reducing immunogenicity or allergenicity of a composition, the method comprising treating the composition to remove cationic proteins, wherein the composition comprises milk or a dairy product and the method comprises removing at least one protein comprising a transcription infidelity peptide sequence selected from SEQ ID NO: 1 to 5. 10 15

18. The method of claim 1, wherein the removed cationic proteins comprise proteins comprising a C-terminal transcription infidelity peptide sequence enriched in basic amino acids resulting from a transcription infidelity deletion. 20

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